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(54) Title: NOVEL NUCLEIC ACIDS AND POLYPEPTIDES

(57) Abstract: The present invention provides novel nucleic acids, novel polypeptide sequences encoded by these nucleic acids and uses thereof.

NOVEL NUCLEIC ACIDS AND POLYPEPTIDES

1. TECHNICAL FIELD

The present invention provides novel polynucleotides and proteins encoded by such polynucleotides, along with uses for these polynucleotides and proteins, for example in therapeutic, diagnostic and research methods.

2. BACKGROUND

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Technology aimed at the discovery of protein factors (including e.g., cytokines, such as lymphokines, interferons, CSFs, chemokines, and interleukins) has matured rapidly over the past decade. The now routine hybridization cloning and expression cloning techniques clone novel polynucleotides "directly" in the sense that they rely on information directly related to the discovered protein (i.e., partial DNA/amino acid sequence of the protein in the case of hybridization cloning; activity of the protein in the case of expression cloning). More recent "indirect" cloning techniques such as signal sequence cloning, which isolates DNA sequences based on the presence of a now well-recognized secretory leader sequence motif, as well as various PCR-based or low stringency hybridization-based cloning techniques, have advanced the state of the art by making available large numbers of DNA/amino acid sequences for proteins that are known to have biological activity, for example, by virtue of their secreted nature in the case of leader sequence cloning, by virtue of their cell or tissue source in the case of PCR-based techniques, or by virtue of structural similarity to other genes of known biological activity.

Identified polynucleotide and polypeptide sequences have numerous applications in, for example, diagnostics, forensics, gene mapping; identification of mutations responsible for genetic disorders or other traits, to assess biodiversity, and to produce many other types of data and products dependent on DNA and amino acid sequences.

3. SUMMARY OF THE INVENTION

The compositions of the present invention include novel isolated polypeptides, novel isolated polynucleotides encoding such polypeptides, including recombinant DNA

molecules, cloned genes or degenerate variants thereof, especially naturally occurring variants such as allelic variants, antisense polynucleotide molecules, and antibodies that specifically recognize one or more epitopes present on such polypeptides, as well as hybridomas producing such antibodies.

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The compositions of the present invention additionally include vectors, including expression vectors, containing the polynucleotides of the invention, cells genetically engineered to contain such polynucleotides and cells genetically engineered to express such polynucleotides.

The present invention relates to a collection or library of at least one novel nucleic acid sequence assembled from expressed sequence tags (ESTs) isolated mainly by sequencing by hybridization (SBH), and in some cases, sequences obtained from one or more public databases. The invention relates also to the proteins encoded by such polynucleotides, along with therapeutic, diagnostic and research utilities for these polynucleotides and proteins. These nucleic acid sequences are designated as SEQ ID NO: 1-438 and are provided in the Sequence Listing. In the nucleic acids provided in the Sequence Listing, A is adenine; C is cytosine; G is guanine; T is thymine; and N is any of the four bases. In the amino acids provided in the Sequence Listing, * corresponds to the stop codon.

The nucleic acid sequences of the present invention also include, nucleic acid sequences that hybridize to the complement of SEQ ID NO: 1 – 438 under stringent hybridization conditions; nucleic acid sequences which are allelic variants or species homologues of any of the nucleic acid sequences recited above, or nucleic acid sequences that encode a peptide comprising a specific domain or truncation of the peptides encoded by SEQ ID NO: 1 – 438. A polynucleotide comprising a nucleotide sequence having at least 90% identity to an identifying sequence of SEQ ID NO: 1 – 438 or a degenerate variant or fragment thereof. The identifying sequence can be 100 base pairs in length.

The nucleic acid sequences of the present invention also include the sequence information from the nucleic acid sequences of SEQ ID NO: 1-438. The sequence information can be a segment of any one of SEQ ID NO: 1-438 that uniquely identifies or represents the sequence information of SEQ ID NO: 1-438.

A collection as used in this application can be a collection of only one polynucleotide. The collection of sequence information or identifying information of each sequence can be provided on a nucleic acid array. In one embodiment, segments of sequence information is provided on a nucleic acid array to detect the polynucleotide that contains the segment. The array can be designed to detect full-match or mismatch to the polynucleotide that contains the segment. The collection can also be provided in a computer-readable format.

This invention also includes the reverse or direct complement of any of the nucleic acid sequences recited above; cloning or expression vectors containing the nucleic acid sequences; and host cells or organisms transformed with these expression vectors. Nucleic acid sequences (or their reverse or direct complements) according to the invention have numerous applications in a variety of techniques known to those skilled in the art of molecular biology, such as use as hybridization probes, use as primers for PCR, use in an array, use in computer-readable media, use in sequencing full-length genes, use for chromosome and gene mapping, use in the recombinant production of protein, and use in the generation of anti-sense DNA or RNA, their chemical analogs and the like.

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In a preferred embodiment, the nucleic acid sequences of SEQ ID NO: 1-438 or novel segments or parts of the nucleic acids of the invention are used as primers in expression assays that are well known in the art. In a particularly preferred embodiment, the nucleic acid sequences of SEQ ID NO: 1-438 or novel segments or parts of the nucleic acids provided herein are used in diagnostics for identifying expressed genes or, as well known in the art and exemplified by Vollrath et al., Science 258:52-59 (1992), as expressed sequence tags for physical mapping of the human genome.

The isolated polynucleotides of the invention include, but are not limited to, a polynucleotide comprising any one of the nucleotide sequences set forth in SEQ ID NO: 1–438; a polynucleotide comprising any of the full length protein coding sequences of SEQ ID NO: 1–438; and a polynucleotide comprising any of the nucleotide sequences of the mature protein coding sequences of SEQ ID NO: 1–438. The polynucleotides of the present invention also include, but are not limited to, a polynucleotide that hybridizes under stringent hybridization conditions to (a) the complement of any one of the nucleotide sequences set forth in SEQ ID NO: 1–438; (b) a nucleotide sequence encoding any one of

the amino acid sequences set forth in the Sequence Listing; (c) a polynucleotide which is an allelic variant of any polynucleotides recited above; (d) a polynucleotide which encodes a species homolog (e.g. orthologs) of any of the proteins recited above; or (e) a polynucleotide that encodes a polypeptide comprising a specific domain or truncation of any of the polypeptides comprising an amino acid sequence set forth in the Sequence Listing.

The isolated polypeptides of the invention include, but are not limited to, a polypeptide comprising any of the amino acid sequences set forth in the Sequence Listing; or the corresponding full length or mature protein. Polypeptides of the invention also include polypeptides with biological activity that are encoded by (a) any of the polynucleotides having a nucleotide sequence set forth in SEQ ID NO: 1-438; or (b) polynucleotides that hybridize to the complement of the polynucleotides of (a) under stringent hybridization conditions. Biologically or immunologically active variants of any of the polypeptide sequences in the Sequence Listing, and "substantial equivalents" thereof (e.g., with at least about 65%, 70%, 75%, 80%, 85%, 90%, 95%, 98% or 99% amino acid sequence identity) that preferably retain biological activity are also contemplated. The polypeptides of the invention may be wholly or partially chemically synthesized but are preferably produced by recombinant means using the genetically engineered cells (e.g. host cells) of the invention.

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The invention also provides compositions comprising a polypeptide of the invention. Polypeptide compositions of the invention may further comprise an acceptable carrier, such as a hydrophilic, e.g., pharmaceutically acceptable, carrier.

The invention also provides host cells transformed or transfected with a polynucleotide of the invention.

The invention also relates to methods for producing a polypeptide of the invention comprising growing a culture of the host cells of the invention in a suitable culture medium under conditions permitting expression of the desired polypeptide, and purifying the polypeptide from the culture or from the host cells. Preferred embodiments include those in which the protein produced by such process is a mature form of the protein.

Polynucleotides according to the invention have numerous applications in a variety of techniques known to those skilled in the art of molecular biology. These techniques include use as hybridization probes, use as oligomers, or primers, for PCR, use for chromosome and gene mapping, use in the recombinant production of protein,

and use in generation of anti-sense DNA or RNA, their chemical analogs and the like. For example, when the expression of an mRNA is largely restricted to a particular cell or tissue type, polynucleotides of the invention can be used as hybridization probes to detect the presence of the particular cell or tissue mRNA in a sample using, e.g., in situ hybridization.

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In other exemplary embodiments, the polynucleotides are used in diagnostics as expressed sequence tags for identifying expressed genes or, as well known in the art and exemplified by Vollrath et al., Science 258:52-59 (1992), as expressed sequence tags for physical mapping of the human genome.

The polypeptides according to the invention can be used in a variety of conventional procedures and methods that are currently applied to other proteins. For example, a polypeptide of the invention can be used to generate an antibody that specifically binds the polypeptide. Such antibodies, particularly monoclonal antibodies, are useful for detecting or quantitating the polypeptide in tissue. The polypeptides of the invention can also be used as molecular weight markers, and as a food supplement.

Methods are also provided for preventing, treating, or ameliorating a medical condition which comprises the step of administering to a mammalian subject a therapeutically effective amount of a composition comprising a polypeptide of the present invention and a pharmaceutically acceptable carrier.

In particular, the polypeptides and polynucleotides of the invention can be utilized, for example, in methods for the prevention and/or treatment of disorders involving aberrant protein expression or biological activity.

The present invention further relates to methods for detecting the presence of the polynucleotides or polypeptides of the invention in a sample. Such methods can, for example, be utilized as part of prognostic and diagnostic evaluation of disorders as recited herein and for the identification of subjects exhibiting a predisposition to such conditions. The invention provides a method for detecting the polynucleotides of the invention in a sample, comprising contacting the sample with a compound that binds to and forms a complex with the polynucleotide of interest for a period sufficient to form the complex and under conditions sufficient to form a complex and detecting the complex such that if a complex is detected, the polynucleotide of interest is detected. The

invention also provides a method for detecting the polypeptides of the invention in a sample comprising contacting the sample with a compound that binds to and forms a complex with the polypeptide under conditions and for a period sufficient to form the complex and detecting the formation of the complex such that if a complex is formed, the polypeptide is detected.

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The invention also provides kits comprising polynucleotide probes and/or monoclonal antibodies, and optionally quantitative standards, for carrying out methods of the invention. Furthermore, the invention provides methods for evaluating the efficacy of drugs, and monitoring the progress of patients, involved in clinical trials for the treatment of disorders as recited above.

The invention also provides methods for the identification of compounds that modulate (i.e., increase or decrease) the expression or activity of the polynucleotides and/or polypeptides of the invention. Such methods can be utilized, for example, for the identification of compounds that can ameliorate symptoms of disorders as recited herein. Such methods can include, but are not limited to, assays for identifying compounds and other substances that interact with (e.g., bind to) the polypeptides of the invention. The invention provides a method for identifying a compound that binds to the polypeptides of the invention comprising contacting the compound with a polypeptide of the invention in a cell for a time sufficient to form a polypeptide/compound complex, wherein the complex drives expression of a reporter gene sequence in the cell; and detecting the complex by detecting the reporter gene sequence expression such that if expression of the reporter gene is detected the compound the binds to a polypeptide of the invention is identified.

The methods of the invention also provides methods for treatment which involve the administration of the polynucleotides or polypeptides of the invention to individuals exhibiting symptoms or tendencies. In addition, the invention encompasses methods for treating diseases or disorders as recited herein comprising administering compounds and other substances that modulate the overall activity of the target gene products.

Compounds and other substances can effect such modulation either on the level of target gene/protein expression or target protein activity.

The polypeptides of the present invention and the polynucleotides encoding them are also useful for the same functions known to one of skill in the art as the polypeptides and polynucleotides to which they have homology (set forth in Table 2); for which they have a signature region (as set forth in Table 3); or for which they have homology to a gene family (as set forth in Table 4). If no homology is set forth for a sequence, then the polypeptides and polynucleotides of the present invention are useful for a variety of applications, as described herein, including use in arrays for detection.

4. DETAILED DESCRIPTION OF THE INVENTION

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4.1 DEFINITIONS

It must be noted that as used herein and in the appended claims, the singular forms "a", "an" and "the" include plural references unless the context clearly dictates otherwise.

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The term "active" refers to those forms of the polypeptide which retain the biologic and/or immunologic activities of any naturally occurring polypeptide. According to the invention, the terms "biologically active" or "biological activity" refer to a protein or peptide having structural, regulatory or biochemical functions of a naturally occurring molecule. Likewise "immunologically active" or "immunological activity" refers to the capability of the natural, recombinant or synthetic polypeptide to induce a specific immune response in appropriate animals or cells and to bind with specific antibodies.

The term "activated cells" as used in this application are those cells which are engaged in extracellular or intracellular membrane trafficking, including the export of secretory or enzymatic molecules as part of a normal or disease process.

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The terms "complementary" or "complementarity" refer to the natural binding of polynucleotides by base pairing. For example, the sequence 5'-AGT-3' binds to the complementary sequence 3'-TCA-5'. Complementarity between two single-stranded molecules may be "partial" such that only some of the nucleic acids bind or it may be "complete" such that total complementarity exists between the single stranded molecules. The degree of complementarity between the nucleic acid strands has significant effects on the efficiency and strength of the hybridization between the nucleic acid strands.

The term "embryonic stem cells (ES)" refers to a cell that can give rise to many differentiated cell types in an embryo or an adult, including the germ cells. The term "germ line stem cells (GSCs)" refers to stem cells derived from primordial stem cells that provide a steady and continuous source of germ cells for the production of gametes. The term "primordial germ cells (PGCs)" refers to a small population of cells set aside from other cell lineages particularly from the yolk sac, mesenteries, or gonadal ridges during embryogenesis that have the potential to differentiate into germ cells and other cells. PGCs are the source from which GSCs and ES cells are derived The PGCs, the GSCs and the ES cells are capable of self-renewal. Thus these cells not only populate the germ line and give rise to a plurality of terminally differentiated cells that comprise the adult specialized organs, but are able to regenerate themselves.

The term "expression modulating fragment," EMF, means a series of nucleotides which modulates the expression of an operably linked ORF or another EMF.

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As used herein, a sequence is said to "modulate the expression of an operably linked sequence" when the expression of the sequence is altered by the presence of the EMF. EMFs include, but are not limited to, promoters, and promoter modulating sequences (inducible elements). One class of EMFs are nucleic acid fragments which induce the expression of an operably linked ORF in response to a specific regulatory factor or physiological event.

The terms "nucleotide sequence" or "nucleic acid" or "polynucleotide" or "oligonculeotide" are used interchangeably and refer to a heteropolymer of nucleotides or the sequence of these nucleotides. These phrases also refer to DNA or RNA of genomic or synthetic origin which may be single-stranded or double-stranded and may represent the sense or the antisense strand, to peptide nucleic acid (PNA) or to any DNA-like or RNA-like material. In the sequences herein A is adenine, C is cytosine, T is thymine, G is guanine and N is A, C, G or T (U). It is contemplated that where the polynucleotide is RNA, the T (thymine) in the sequences provided herein is substituted with U (uracil). Generally, nucleic acid segments provided by this invention may be assembled from fragments of the genome and short oligonucleotide linkers, or from a series of oligonucleotides, or from individual nucleotides, to provide a synthetic nucleic acid

which is capable of being expressed in a recombinant transcriptional unit comprising regulatory elements derived from a microbial or viral operon, or a eukaryotic gene.

The terms "oligonucleotide fragment" or a "polynucleotide fragment", "portion," or "segment" or "probe" or "primer" are used interchangeably and refer to a sequence of nucleotide residues which are at least about 5 nucleotides, more preferably at least about 7 nucleotides, more preferably at least about 9 nucleotides, more preferably at least about 11 nucleotides and most preferably at least about 17 nucleotides. The fragment is preferably less than about 500 nucleotides, preferably less than about 200 nucleotides, more preferably less than about 100 nucleotides, more preferably less than about 50 nucleotides and most preferably less than 30 nucleotides. Preferably the probe is from about 6 nucleotides to about 200 nucleotides, preferably from about 15 to about 50 nucleotides, more preferably from about 17 to 30 nucleotides and most preferably from about 20 to 25 nucleotides. Preferably the fragments can be used in polymerase chain reaction (PCR), various hybridization procedures or microarray procedures to identify or amplify identical or related parts of mRNA or DNA molecules. A fragment or segment may uniquely identify each polynucleotide sequence of the present invention. Preferably the fragment comprises a sequence substantially similar to any one of SEQ ID NOs:1-438.

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Probes may, for example, be used to determine whether specific mRNA molecules are present in a cell or tissue or to isolate similar nucleic acid sequences from chromosomal DNA as described by Walsh et al. (Walsh, P.S. et al., 1992, PCR Methods Appl 1:241-250). They may be labeled by nick translation, Klenow fill-in reaction, PCR, or other methods well known in the art. Probes of the present invention, their preparation and/or labeling are elaborated in Sambrook, J. et al., 1989, Molecular Cloning: A Laboratory Manual, Cold Spring Harbor Laboratory, NY; or Ausubel, F.M. et al., 1989, Current Protocols in Molecular Biology, John Wiley & Sons, New York NY, both of which are incorporated herein by reference in their entirety.

The nucleic acid sequences of the present invention also include the sequence information from the nucleic acid sequences of SEQ ID NOs: 1-438. The sequence information can be a segment of any one of SEQ ID NOs: 1-438 that uniquely identifies or represents the sequence information of that sequence of SEQ ID NO: 1-438. One such

segment can be a twenty-mer nucleic acid sequence because the probability that a twenty-mer is fully matched in the human genome is 1 in 300. In the human genome, there are three billion base pairs in one set of chromosomes. Because 4^{20} possible twenty-mers exist, there are 300 times more twenty-mers than there are base pairs in a set of human chromosomes. Using the same analysis, the probability for a seventeen-mer to be fully matched in the human genome is approximately 1 in 5. When these segments are used in arrays for expression studies, fifteen-mer segments can be used. The probability that the fifteen-mer is fully matched in the expressed sequences is also approximately one in five because expressed sequences comprise less than approximately 5% of the entire genome sequence.

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Similarly, when using sequence information for detecting a single mismatch, a segment can be a twenty-five mer. The probability that the twenty-five mer would appear in a human genome with a single mismatch is calculated by multiplying the probability for a full match $(1 \div 4^{25})$ times the increased probability for mismatch at each nucleotide position (3×25) . The probability that an eighteen mer with a single mismatch can be detected in an array for expression studies is approximately one in five. The probability that a twenty-mer with a single mismatch can be detected in a human genome is approximately one in five.

The term "open reading frame," ORF, means a series of nucleotide triplets coding for amino acids without any termination codons and is a sequence translatable into protein.

The terms "operably linked" or "operably associated" refer to functionally related nucleic acid sequences. For example, a promoter is operably associated or operably linked with a coding sequence if the promoter controls the transcription of the coding sequence. While operably linked nucleic acid sequences can be contiguous and in the same reading frame, certain genetic elements e.g. repressor genes are not contiguously linked to the coding sequence but still control transcription/translation of the coding sequence.

The term "pluripotent" refers to the capability of a cell to differentiate into a number of differentiated cell types that are present in an adult organism. A pluripotent cell is restricted in its differentiation capability in comparison to a totipotent cell.

The terms "polypeptide" or "peptide" or "amino acid sequence" refer to an oligopeptide, peptide, polypeptide or protein sequence or fragment thereof and to naturally occurring or synthetic molecules. A polypeptide "fragment," "portion," or "segment" is a stretch of amino acid residues of at least about 5 amino acids, preferably at least about 7 amino acids, more preferably at least about 9 amino acids and most preferably at least about 17 or more amino acids. The peptide preferably is not greater than about 200 amino acids, more preferably less than 150 amino acids and most preferably less than 100 amino acids. Preferably the peptide is from about 5 to about 200 amino acids. To be active, any polypeptide must have sufficient length to display biological and/or immunological activity.

The term "naturally occurring polypeptide" refers to polypeptides produced by cells that have not been genetically engineered and specifically contemplates various polypeptides arising from post-translational modifications of the polypeptide including, but not limited to, acetylation, carboxylation, glycosylation, phosphorylation, lipidation and acylation.

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The term "translated protein coding portion" means a sequence which encodes for the full length protein which may include any leader sequence or any processing sequence.

The term "mature protein coding sequence" means a sequence which encodes a peptide or protein without a signal or leader sequence. The "mature protein portion" means that portion of the protein which does not include a signal or leader sequence. The peptide may have been produced by processing in the cell which removes any leader/signal sequence. The mature protein portion may or may not include the initial methionine residue. The methionine residue may be removed from the protein during processing in the cell. The peptide may be produced synthetically or the protein may have been produced using a polynucleotide only encoding for the mature protein coding sequence.

The term "derivative" refers to polypeptides chemically modified by such techniques as ubiquitination, labeling (e.g., with radionuclides or various enzymes), covalent polymer attachment such as pegylation (derivatization with polyethylene glycol)

and insertion or substitution by chemical synthesis of amino acids such as ornithine, which do not normally occur in human proteins.

The term "variant" (or "analog") refers to any polypeptide differing from naturally occurring polypeptides by amino acid insertions, deletions, and substitutions, created using, e.g., recombinant DNA techniques. Guidance in determining which amino acid residues may be replaced, added or deleted without abolishing activities of interest, may be found by comparing the sequence of the particular polypeptide with that of homologous peptides and minimizing the number of amino acid sequence changes made in regions of high homology (conserved regions) or by replacing amino acids with consensus sequence.

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Alternatively, recombinant variants encoding these same or similar polypeptides may be synthesized or selected by making use of the "redundancy" in the genetic code. Various codon substitutions, such as the silent changes which produce various restriction sites, may be introduced to optimize cloning into a plasmid or viral vector or expression in a particular prokaryotic or eukaryotic system. Mutations in the polynucleotide sequence may be reflected in the polypeptide or domains of other peptides added to the polypeptide to modify the properties of any part of the polypeptide, to change characteristics such as ligand-binding affinities, interchain affinities, or degradation/turnover rate.

Preferably, amino acid "substitutions" are the result of replacing one amino acid with another amino acid having similar structural and/or chemical properties, i.e., conservative amino acid replacements. "Conservative" amino acid substitutions may be made on the basis of similarity in polarity, charge, solubility, hydrophobicity, hydrophobicity, and/or the amphipathic nature of the residues involved. For example, nonpolar (hydrophobic) amino acids include alanine, leucine, isoleucine, valine, proline, phenylalanine, tryptophan, and methionine; polar neutral amino acids include glycine, serine, threonine, cysteine, tyrosine, asparagine, and glutamine; positively charged (basic) amino acids include arginine, lysine, and histidine; and negatively charged (acidic) amino acids include aspartic acid and glutamic acid. "Insertions" or "deletions" are preferably in the range of about 1 to 20 amino acids, more preferably 1 to 10 amino acids. The variation allowed may be experimentally determined by systematically making insertions,

deletions, or substitutions of amino acids in a polypeptide molecule using recombinant DNA techniques and assaying the resulting recombinant variants for activity.

Alternatively, where alteration of function is desired, insertions, deletions or non-conservative alterations can be engineered to produce altered polypeptides. Such alterations can, for example, alter one or more of the biological functions or biochemical characteristics of the polypeptides of the invention. For example, such alterations may change polypeptide characteristics such as ligand-binding affinities, interchain affinities, or degradation/turnover rate. Further, such alterations can be selected so as to generate polypeptides that are better suited for expression, scale up and the like in the host cells chosen for expression. For example, cysteine residues can be deleted or substituted with another amino acid residue in order to eliminate disulfide bridges.

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The terms "purified" or "substantially purified" as used herein denotes that the indicated nucleic acid or polypeptide is present in the substantial absence of other biological macromolecules, e.g., polynucleotides, proteins, and the like. In one embodiment, the polynucleotide or polypeptide is purified such that it constitutes at least 95% by weight, more preferably at least 99% by weight, of the indicated biological macromolecules present (but water, buffers, and other small molecules, especially molecules having a molecular weight of less than 1000 daltons, can be present).

The term "isolated" as used herein refers to a nucleic acid or polypeptide separated from at least one other component (e.g., nucleic acid or polypeptide) present with the nucleic acid or polypeptide in its natural source. In one embodiment, the nucleic acid or polypeptide is found in the presence of (if anything) only a solvent, buffer, ion, or other component normally present in a solution of the same. The terms "isolated" and "purified" do not encompass nucleic acids or polypeptides present in their natural source.

The term "recombinant," when used herein to refer to a polypeptide or protein, means that a polypeptide or protein is derived from recombinant (e.g., microbial, insect, or mammalian) expression systems. "Microbial" refers to recombinant polypeptides or proteins made in bacterial or fungal (e.g., yeast) expression systems. As a product, "recombinant microbial" defines a polypeptide or protein essentially free of native endogenous substances and unaccompanied by associated native glycosylation. Polypeptides or proteins expressed in most bacterial cultures, e.g., E. coli, will be free of

glycosylation modifications; polypeptides or proteins expressed in yeast will have a glycosylation pattern in general different from those expressed in mammalian cells.

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The term "recombinant expression vehicle or vector" refers to a plasmid or phage or virus or vector, for expressing a polypeptide from a DNA (RNA) sequence. An expression vehicle can comprise a transcriptional unit comprising an assembly of (1) a genetic element or elements having a regulatory role in gene expression, for example, promoters or enhancers, (2) a structural or coding sequence which is transcribed into mRNA and translated into protein, and (3) appropriate transcription initiation and termination sequences. Structural units intended for use in yeast or eukaryotic expression systems preferably include a leader sequence enabling extracellular secretion of translated protein by a host cell. Alternatively, where recombinant protein is expressed without a leader or transport sequence, it may include an amino terminal methionine residue. This residue may or may not be subsequently cleaved from the expressed recombinant protein to provide a final product.

The term "recombinant expression system" means host cells which have stably integrated a recombinant transcriptional unit into chromosomal DNA or carry the recombinant transcriptional unit extrachromosomally. Recombinant expression systems as defined herein will express heterologous polypeptides or proteins upon induction of the regulatory elements linked to the DNA segment or synthetic gene to be expressed. This term also means host cells which have stably integrated a recombinant genetic element or elements having a regulatory role in gene expression, for example, promoters or enhancers. Recombinant expression systems as defined herein will express polypeptides or proteins endogenous to the cell upon induction of the regulatory elements linked to the endogenous DNA segment or gene to be expressed. The cells can be prokaryotic or eukaryotic.

The term "secreted" includes a protein that is transported across or through a membrane, including transport as a result of signal sequences in its amino acid sequence when it is expressed in a suitable host cell. "Secreted" proteins include without limitation proteins secreted wholly (e.g., soluble proteins) or partially (e.g., receptors) from the cell in which they are expressed. "Secreted" proteins also include without limitation proteins that are transported across the membrane of the endoplasmic reticulum. "Secreted"

proteins are also intended to include proteins containing non-typical signal sequences (e.g. Interleukin-1 Beta, see Krasney, P.A. and Young, P.R. (1992) Cytokine 4(2):134-143) and factors released from damaged cells (e.g. Interleukin-1 Receptor Antagonist, see Arend, W.P. et. al. (1998) Annu. Rev. Immunol. 16:27-55)

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Where desired, an expression vector may be designed to contain a "signal or leader sequence" which will direct the polypeptide through the membrane of a cell. Such a sequence may be naturally present on the polypeptides of the present invention or provided from heterologous protein sources by recombinant DNA techniques.

The term "stringent" is used to refer to conditions that are commonly understood in the art as stringent. Stringent conditions can include highly stringent conditions (i.e., hybridization to filter-bound DNA in 0.5 M NaHPO₄, 7% sodium dodecyl sulfate (SDS), 1 mM EDTA at 65°C, and washing in 0.1X SSC/0.1% SDS at 68°C), and moderately stringent conditions (i.e., washing in 0.2X SSC/0.1% SDS at 42°C). Other exemplary hybridization conditions are described herein in the examples.

In instances of hybridization of deoxyoligonucleotides, additional exemplary stringent hybridization conditions include washing in 6X SSC/0.05% sodium pyrophosphate at 37°C (for 14-base oligonucleotides), 48°C (for 17-base oligos), 55°C (for 20-base oligonucleotides), and 60°C (for 23-base oligonucleotides).

As used herein, "substantially equivalent" or "substantially similar" can refer both to nucleotide and amino acid sequences, for example a mutant sequence, that varies from a reference sequence by one or more substitutions, deletions, or additions, the net effect of which does not result in an adverse functional dissimilarity between the reference and subject sequences. Typically, such a substantially equivalent sequence varies from one of those listed herein by no more than about 35% (i.e., the number of individual residue substitutions, additions, and/or deletions in a substantially equivalent sequence, as compared to the corresponding reference sequence, divided by the total number of residues in the substantially equivalent sequence is about 0.35 or less). Such a sequence is said to have 65% sequence identity to the listed sequence. In one embodiment, a substantially equivalent, e.g., mutant, sequence of the invention varies from a listed sequence by no more than 30% (70% sequence identity); in a variation of this embodiment, by no more than 25% (75% sequence identity); and in a further variation of

this embodiment, by no more than 20% (80% sequence identity) and in a further variation of this embodiment, by no more than 10% (90% sequence identity) and in a further variation of this embodiment, by no more that 5% (95% sequence identity). Substantially equivalent, e.g., mutant, amino acid sequences according to the invention preferably have at least 80% sequence identity with a listed amino acid sequence, more preferably at least 85% sequence identity, more preferably at least 90% sequence identity, more preferably at least 95% sequence identity, more preferably at least 98% sequence identity, and most preferably at least 99% sequence identity. Substantially equivalent nucleotide sequence of the invention can have lower percent sequence identities, taking into account, for example, the redundancy or degeneracy of the genetic code. Preferably, the nucleotide sequence has at least about 65% identity, more preferably at least about 75% identity, more preferably at least about 80% sequence identity, more preferably at least 85% sequence identity, more preferably at least 90% sequence identity, more preferably at least about 95% sequence identity, more preferably at least 98% sequence identity, and most preferably at least 99% sequence identity. For the purposes of the present invention, sequences having substantially equivalent biological activity and substantially equivalent expression characteristics are considered substantially equivalent. For the purposes of determining equivalence, truncation of the mature sequence (e.g., via a mutation which creates a spurious stop codon) should be disregarded. Sequence identity may be determined, e.g., using the Jotun Hein method (Hein, J. (1990) Methods Enzymol. 183:626-645). Identity between sequences can also be determined by other methods known in the art, e.g. by varying hybridization conditions.

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The term "totipotent" refers to the capability of a cell to differentiate into all of the cell types of an adult organism.

The term "transformation" means introducing DNA into a suitable host cell so that the DNA is replicable, either as an extrachromosomal element, or by chromosomal integration. The term "transfection" refers to the taking up of an expression vector by a suitable host cell, whether or not any coding sequences are in fact expressed. The term "infection" refers to the introduction of nucleic acids into a suitable host cell by use of a virus or viral vector.

As used herein, an "uptake modulating fragment," UMF, means a series of nucleotides which mediate the uptake of a linked DNA fragment into a cell. UMFs can be readily identified using known UMFs as a target sequence or target motif with the computer-based systems described below. The presence and activity of a UMF can be confirmed by attaching the suspected UMF to a marker sequence. The resulting nucleic acid molecule is then incubated with an appropriate host under appropriate conditions and the uptake of the marker sequence is determined. As described above, a UMF will increase the frequency of uptake of a linked marker sequence.

Each of the above terms is meant to encompass all that is described for each, unless the context dictates otherwise.

4.2 NUCLEIC ACIDS OF THE INVENTION

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Nucleotide sequences of the invention are set forth in the Sequence Listing. The isolated polynucleotides of the invention include a polynucleotide comprising the nucleotide sequences of SEQ ID NO: 1 - 438; a polynucleotide encoding any one of the peptide sequences of SEQ ID NO:1 – 438; and a polynucleotide comprising the nucleotide sequence encoding the mature protein coding sequence of the polynucleotides of any one of SEQ ID NO: 1 - 438. The polynucleotides of the present invention also include, but are not limited to, a polynucleotide that hybridizes under stringent conditions to (a) the complement of any of the nucleotides sequences of SEQ ID NO: 1-438; (b) nucleotide sequences encoding any one of the amino acid sequences set forth in the Sequence Listing; (c) a polynucleotide which is an allelic variant of any polynucleotide recited above; (d) a polynucleotide which encodes a species homolog of any of the proteins recited above; or (e) a polynucleotide that encodes a polypeptide comprising a specific domain or truncation of the polypeptides of SEQ ID NO: 1-438. Domains of interest may depend on the nature of the encoded polypeptide; e.g., domains in receptorlike polypeptides include ligand-binding, extracellular, transmembrane, or cytoplasmic domains, or combinations thereof; domains in immunoglobulin-like proteins include the variable immunoglobulin-like domains; domains in enzyme-like polypeptides include catalytic and substrate binding domains; and domains in ligand polypeptides include receptor-binding domains.

The polynucleotides of the invention include naturally occurring or wholly or partially synthetic DNA, e.g., cDNA and genomic DNA, and RNA, e.g., mRNA. The polynucleotides may include all of the coding region of the cDNA or may represent a portion of the coding region of the cDNA.

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The present invention also provides genes corresponding to the cDNA sequences disclosed herein. The corresponding genes can be isolated in accordance with known methods using the sequence information disclosed herein. Such methods include the preparation of probes or primers from the disclosed sequence information for identification and/or amplification of genes in appropriate genomic libraries or other sources of genomic materials. Further 5' and 3' sequence can be obtained using methods known in the art. For example, full length cDNA or genomic DNA that corresponds to any of the polynucleotides of SEQ ID NO: 1 – 438 can be obtained by screening appropriate cDNA or genomic DNA libraries under suitable hybridization conditions using any of the polynucleotides of SEQ ID NO: 1 – 438 or a portion thereof as a probe. Alternatively, the polynucleotides of SEQ ID NO: 1 – 438 may be used as the basis for suitable primer(s) that allow identification and/or amplification of genes in appropriate genomic DNA or cDNA libraries.

The nucleic acid sequences of the invention can be assembled from ESTs and sequences (including cDNA and genomic sequences) obtained from one or more public databases, such as dbEST, gbpri, and UniGene. The EST sequences can provide identifying sequence information, representative fragment or segment information, or novel segment information for the full-length gene.

The polynucleotides of the invention also provide polynucleotides including nucleotide sequences that are substantially equivalent to the polynucleotides recited above. Polynucleotides according to the invention can have, e.g., at least about 65%, at least about 70%, at least about 75%, at least about 80%, 81%, 82%, 83%, 84%, more typically at least about 85%, 86%, 87%, 88%, 89%, more typically at least about 90%, 91%, 92%, 93%, 94%, and even more typically at least about 95%, 96%, 97%, 98%, 99% sequence identity to a polynucleotide recited above.

Included within the scope of the nucleic acid sequences of the invention are nucleic acid sequence fragments that hybridize under stringent conditions to any of the nucleotide sequences of SEQ ID NO: 1 - 438, or complements thereof, which fragment is

greater than about 5 nucleotides, preferably 7 nucleotides, more preferably greater than 9 nucleotides and most preferably greater than 17 nucleotides. Fragments of, e.g. 15, 17, or 20 nucleotides or more that are selective for (i.e. specifically hybridize to any one of the polynucleotides of the invention) are contemplated. Probes capable of specifically hybridizing to a polynucleotide can differentiate polynucleotide sequences of the invention from other polynucleotide sequences in the same family of genes or can differentiate human genes from genes of other species, and are preferably based on unique nucleotide sequences.

The sequences falling within the scope of the present invention are not limited to these specific sequences, but also include allelic and species variations thereof. Allelic and species variations can be routinely determined by comparing the sequence provided in SEQ ID NO: 1 - 438, a representative fragment thereof, or a nucleotide sequence at least 90% identical, preferably 95% identical, to SEQ ID NOs: 1 - 438 with a sequence from another isolate of the same species. Furthermore, to accommodate codon variability, the invention includes nucleic acid molecules coding for the same amino acid sequences as do the specific ORFs disclosed herein. In other words, in the coding region of an ORF, substitution of one codon for another codon that encodes the same amino acid is expressly contemplated.

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The nearest neighbor or homology result for the nucleic acids of the present invention, including SEQ ID NOs: 1 - 438, can be obtained by searching a database using an algorithm or a program. Preferably, a BLAST which stands for Basic Local Alignment Search Tool is used to search for local sequence alignments (Altshul, S.F. J Mol. Evol. 36 290-300 (1993) and Altschul S.F. et al. J. Mol. Biol. 21:403-410 (1990)). Alternatively a FASTA version 3 search against Genpept, using Fastxy algorithm.

Species homologs (or orthologs) of the disclosed polynucleotides and proteins are also provided by the present invention. Species homologs may be isolated and identified by making suitable probes or primers from the sequences provided herein and screening a suitable nucleic acid source from the desired species.

The invention also encompasses allelic variants of the disclosed polynucleotides or proteins; that is, naturally-occurring alternative forms of the isolated polynucleotide which also encode proteins which are identical, homologous or related to that encoded by the polynucleotides.

The nucleic acid sequences of the invention are further directed to sequences which encode variants of the described nucleic acids. These amino acid sequence variants may be prepared by methods known in the art by introducing appropriate nucleotide changes into a native or variant polynucleotide. There are two variables in the construction of amino acid sequence variants: the location of the mutation and the nature of the mutation. Nucleic acids encoding the amino acid sequence variants are preferably constructed by mutating the polynucleotide to encode an amino acid sequence that does not occur in nature. These nucleic acid alterations can be made at sites that differ in the nucleic acids from different species (variable positions) or in highly conserved regions (constant regions). Sites at such locations will typically be modified in series, e.g., by substituting first with conservative choices (e.g., hydrophobic amino acid to a different hydrophobic amino acid) and then with more distant choices (e.g., hydrophobic amino acid to a charged amino acid), and then deletions or insertions may be made at the target site. Amino acid sequence deletions generally range from about 1 to 30 residues, preferably about 1 to 10 residues, and are typically contiguous. Amino acid insertions include amino- and/or carboxyl-terminal fusions ranging in length from one to one hundred or more residues, as well as intrasequence insertions of single or multiple amino acid residues. Intrasequence insertions may range generally from about 1 to 10 amino residues, preferably from 1 to 5 residues. Examples of terminal insertions include the heterologous signal sequences necessary for secretion or for intracellular targeting in different host cells and sequences such as FLAG or poly-histidine sequences useful for purifying the expressed protein.

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In a preferred method, polynucleotides encoding the novel amino acid sequences are changed via site-directed mutagenesis. This method uses oligonucleotide sequences to alter a polynucleotide to encode the desired amino acid variant, as well as sufficient adjacent nucleotides on both sides of the changed amino acid to form a stable duplex on either side of the site of being changed. In general, the techniques of site-directed mutagenesis are well known to those of skill in the art and this technique is exemplified by publications such as, Edelman et al., *DNA* 2:183 (1983). A versatile and efficient method for producing site-specific changes in a polynucleotide sequence was published by Zoller and Smith, *Nucleic Acids Res.* 10:6487-6500 (1982). PCR may also be used to

create amino acid sequence variants of the novel nucleic acids. When small amounts of template DNA are used as starting material, primer(s) that differs slightly in sequence from the corresponding region in the template DNA can generate the desired amino acid variant. PCR amplification results in a population of product DNA fragments that differ from the polynucleotide template encoding the polypeptide at the position specified by the primer. The product DNA fragments replace the corresponding region in the plasmid and this gives a polynucleotide encoding the desired amino acid variant.

A further technique for generating amino acid variants is the cassette mutagenesis technique described in Wells et al., *Gene* 34:315 (1985); and other mutagenesis techniques well known in the art, such as, for example, the techniques in Sambrook et al., supra, and *Current Protocols in Molecular Biology*, Ausubel et al. Due to the inherent degeneracy of the genetic code, other DNA sequences which encode substantially the same or a functionally equivalent amino acid sequence may be used in the practice of the invention for the cloning and expression of these novel nucleic acids. Such DNA sequences include those which are capable of hybridizing to the appropriate novel nucleic acid sequence under stringent conditions.

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Polynucleotides encoding preferred polypeptide truncations of the invention can be used to generate polynucleotides encoding chimeric or fusion proteins comprising one or more domains of the invention and heterologous protein sequences.

The polynucleotides of the invention additionally include the complement of any of the polynucleotides recited above. The polynucleotide can be DNA (genomic, cDNA, amplified, or synthetic) or RNA. Methods and algorithms for obtaining such polynucleotides are well known to those of skill in the art and can include, for example, methods for determining hybridization conditions that can routinely isolate polynucleotides of the desired sequence identities.

In accordance with the invention, polynucleotide sequences comprising the mature protein coding sequences corresponding to any one of SEQ ID NO: 1-438, or functional equivalents thereof, may be used to generate recombinant DNA molecules that direct the expression of that nucleic acid, or a functional equivalent thereof, in appropriate host cells. Also included are the cDNA inserts of any of the clones identified herein.

A polynucleotide according to the invention can be joined to any of a variety of other nucleotide sequences by well-established recombinant DNA techniques (see Sambrook J et al. (1989) Molecular Cloning: A Laboratory Manual, Cold Spring Harbor Laboratory, NY). Useful nucleotide sequences for joining to polynucleotides include an assortment of vectors, e.g., plasmids, cosmids, lambda phage derivatives, phagemids, and the like, that are well known in the art. Accordingly, the invention also provides a vector including a polynucleotide of the invention and a host cell containing the polynucleotide. In general, the vector contains an origin of replication functional in at least one organism, convenient restriction endonuclease sites, and a selectable marker for the host cell. Vectors according to the invention include expression vectors, replication vectors, probe generation vectors, and sequencing vectors. A host cell according to the invention can be a prokaryotic or eukaryotic cell and can be a unicellular organism or part of a multicellular organism.

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The present invention further provides recombinant constructs comprising a nucleic acid having any of the nucleotide sequences of SEQ ID NOs: 1 - 438 or a fragment thereof or any other polynucleotides of the invention. In one embodiment, the recombinant constructs of the present invention comprise a vector, such as a plasmid or viral vector, into which a nucleic acid having any of the nucleotide sequences of SEQ ID NOs: 1 - 438 or a fragment thereof is inserted, in a forward or reverse orientation. In the case of a vector comprising one of the ORFs of the present invention, the vector may further comprise regulatory sequences, including for example, a promoter, operably linked to the ORF. Large numbers of suitable vectors and promoters are known to those of skill in the art and are commercially available for generating the recombinant constructs of the present invention. The following vectors are provided by way of example. Bacterial: pBs, phagescript, PsiX174, pBluescript SK, pBs KS, pNH8a, pNH16a, pNH18a, pNH46a (Stratagene); pTrc99A, pKK223-3, pKK233-3, pDR540, pRIT5 (Pharmacia). Eukaryotic: pWLneo, pSV2cat, pOG44, PXTI, pSG (Stratagene) pSVK3, pBPV, pMSG, pSVL (Pharmacia).

The isolated polynucleotide of the invention may be operably linked to an expression control sequence such as the pMT2 or pED expression vectors disclosed in Kaufman et al., *Nucleic Acids Res.* 19, 4485-4490 (1991), in order to produce the protein

recombinantly. Many suitable expression control sequences are known in the art. General methods of expressing recombinant proteins are also known and are exemplified in R. Kaufman, *Methods in Enzymology* 185, 537-566 (1990). As defined herein "operably linked" means that the isolated polynucleotide of the invention and an expression control sequence are situated within a vector or cell in such a way that the protein is expressed by a host cell which has been transformed (transfected) with the ligated polynucleotide/expression control sequence.

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Promoter regions can be selected from any desired gene using CAT (chloramphenicol transferase) vectors or other vectors with selectable markers. Two appropriate vectors are pKK232-8 and pCM7. Particular named bacterial promoters include lacI, lacZ, T3, T7, gpt, lambda PR, and trc. Eukaryotic promoters include CMV immediate early, HSV thymidine kinase, early and late SV40, LTRs from retrovirus, and mouse metallothionein-I. Selection of the appropriate vector and promoter is well within the level of ordinary skill in the art. Generally, recombinant expression vectors will include origins of replication and selectable markers permitting transformation of the host cell, e.g., the ampicillin resistance gene of E. coli and S. cerevisiae TRP1 gene, and a promoter derived from a highly-expressed gene to direct transcription of a downstream structural sequence. Such promoters can be derived from operons encoding glycolytic enzymes such as 3-phosphoglycerate kinase (PGK), a-factor, acid phosphatase, or heat shock proteins, among others. The heterologous structural sequence is assembled in appropriate phase with translation initiation and termination sequences, and preferably, a leader sequence capable of directing secretion of translated protein into the periplasmic space or extracellular medium. Optionally, the heterologous sequence can encode a fusion protein including an amino terminal identification peptide imparting desired characteristics, e.g., stabilization or simplified purification of expressed recombinant product. Useful expression vectors for bacterial use are constructed by inserting a structural DNA sequence encoding a desired protein together with suitable translation initiation and termination signals in operable reading phase with a functional promoter. The vector will comprise one or more phenotypic selectable markers and an origin of replication to ensure maintenance of the vector and to, if desirable, provide amplification within the host. Suitable prokaryotic hosts for transformation include E. coli, Bacillus

subtilis, Salmonella typhimurium and various species within the genera Pseudomonas, Streptomyces, and Staphylococcus, although others may also be employed as a matter of choice.

As a representative but non-limiting example, useful expression vectors for bacterial use can comprise a selectable marker and bacterial origin of replication derived from commercially available plasmids comprising genetic elements of the well known cloning vector pBR322 (ATCC 37017). Such commercial vectors include, for example, pKK223-3 (Pharmacia Fine Chemicals, Uppsala, Sweden) and GEM 1 (Promega Biotech, Madison, WI, USA). These pBR322 "backbone" sections are combined with an appropriate promoter and the structural sequence to be expressed. Following transformation of a suitable host strain and growth of the host strain to an appropriate cell density, the selected promoter is induced or derepressed by appropriate means (e.g., temperature shift or chemical induction) and cells are cultured for an additional period. Cells are typically harvested by centrifugation, disrupted by physical or chemical means, and the resulting crude extract retained for further purification.

Polynucleotides of the invention can also be used to induce immune responses. For example, as described in Fan et al., *Nat. Biotech.* 17:870-872 (1999), incorporated herein by reference, nucleic acid sequences encoding a polypeptide may be used to generate antibodies against the encoded polypeptide following topical administration of naked plasmid DNA or following injection, and preferably intra-muscular injection of the DNA. The nucleic acid sequences are preferably inserted in a recombinant expression vector and may be in the form of naked DNA.

4.3 ANTISENSE

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Another aspect of the invention pertains to isolated antisense nucleic acid molecules that are hybridizable to or complementary to the nucleic acid molecule comprising the nucleotide sequence of SEQ ID NO: 1 - 438, or fragments, analogs or derivatives thereof. An "antisense" nucleic acid comprises a nucleotide sequence that is complementary to a "sense" nucleic acid encoding a protein, e.g., complementary to the coding strand of a double-stranded cDNA molecule or complementary to an mRNA sequence. In specific aspects, antisense nucleic acid molecules are provided that

comprise a sequence complementary to at least about 10, 25, 50, 100, 250 or 500 nucleotides or an entire coding strand, or to only a portion thereof. Nucleic acid molecules encoding fragments, homologs, derivatives and analogs of a protein of any of SEQ ID NO: 1 - 438 or antisense nucleic acids complementary to a nucleic acid sequence of SEQ ID NO: 1 - 438 are additionally provided.

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In one embodiment, an antisense nucleic acid molecule is antisense to a "coding region" of the coding strand of a nucleotide sequence of the invention. The term "coding region" refers to the region of the nucleotide sequence comprising codons which are translated into amino acid residues. In another embodiment, the antisense nucleic acid molecule is antisense to a "noncoding region" of the coding strand of a nucleotide sequence of the invention. The term "noncoding region" refers to 5' and 3' sequences that flank the coding region that are not translated into amino acids (i.e., also referred to as 5' and 3' untranslated regions).

Given the coding strand sequences encoding a nucleic acid disclosed herein (e.g., SEQ ID NO: 1 - 438, antisense nucleic acids of the invention can be designed according to the rules of Watson and Crick or Hoogsteen base pairing. The antisense nucleic acid molecule can be complementary to the entire coding region of an mRNA, but more preferably is an oligonucleotide that is antisense to only a portion of the coding or noncoding region of an mRNA. For example, the antisense oligonucleotide can be complementary to the region surrounding the translation start site of an mRNA. An antisense oligonucleotide can be, for example, about 5, 10, 15, 20, 25, 30, 35, 40, 45 or 50 nucleotides in length. An antisense nucleic acid of the invention can be constructed using chemical synthesis or enzymatic ligation reactions using procedures known in the art. For example, an antisense nucleic acid (e.g., an antisense oligonucleotide) can be chemically synthesized using naturally occurring nucleotides or variously modified nucleotides designed to increase the biological stability of the molecules or to increase the physical stability of the duplex formed between the antisense and sense nucleic acids, e.g., phosphorothioate derivatives and acridine substituted nucleotides can be used.

Examples of modified nucleotides that can be used to generate the antisense nucleic acid include: 5-fluorouracil, 5-bromouracil, 5-chlorouracil, 5-iodouracil, hypoxanthine, xanthine, 4-acetylcytosine, 5-(carboxyhydroxylmethyl) uracil,

5-carboxymethylaminomethyl-2-thiouridine, 5-carboxymethylaminomethyluracil, dihydrouracil, beta-D-galactosylqueosine, inosine, N6-isopentenyladenine, 1-methylguanine, 1-methylguanine, 2,2-dimethylguanine, 2-methylguanine, 2-methylguanine, 3-methylcytosine, 5-methylcytosine, N6-adenine, 7-methylguanine, 5-methylguanine, 5-methylaminomethyldracil, 5-methoxyaminomethyl-2-thiouracil, beta-D-mannosylqueosine, 5'-methoxycarboxymethyluracil, 5-methoxyuracil, 2-methylthio-N6-isopentenyladenine, uracil-5-oxyacetic acid (v), wybutoxosine, pseudouracil, queosine, 2-thiocytosine, 5-methyl-2-thiouracil, 2-thiouracil, 4-thiouracil, 5-methyluracil, uracil-5-oxyacetic acid methylester, uracil-5-oxyacetic acid (v), 5-methyl-2-thiouracil, 3-(3-amino-3-N-2-carboxypropyl) uracil, (acp3)w, and 2,6-diaminopurine. Alternatively, the antisense nucleic acid can be produced biologically using an expression vector into which a nucleic acid has been subcloned in an antisense orientation (i.e., RNA transcribed from the inserted nucleic acid will be of an antisense orientation to a target nucleic acid of interest, described further in the following

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subsection).

The antisense nucleic acid molecules of the invention are typically administered to a subject or generated in situ such that they hybridize with or bind to cellular mRNA and/or genomic DNA encoding a protein according to the invention to thereby inhibit expression of the protein, e.g., by inhibiting transcription and/or translation. The 20 hybridization can be by conventional nucleotide complementarity to form a stable duplex, or, for example, in the case of an antisense nucleic acid molecule that binds to DNA duplexes, through specific interactions in the major groove of the double helix. An example of a route of administration of antisense nucleic acid molecules of the invention includes direct injection at a tissue site. Alternatively, antisense nucleic acid molecules 25 can be modified to target selected cells and then administered systemically. For example, for systemic administration, antisense molecules can be modified such that they specifically bind to receptors or antigens expressed on a selected cell surface, e.g., by linking the antisense nucleic acid molecules to peptides or antibodies that bind to cell surface receptors or antigens. The antisense nucleic acid molecules can also be delivered 30 to cells using the vectors described herein. To achieve sufficient intracellular concentrations of antisense molecules, vector constructs in which the antisense nucleic

acid molecule is placed under the control of a strong pol II or pol III promoter are preferred.

In yet another embodiment, the antisense nucleic acid molecule of the invention is an α-anomeric nucleic acid molecule. An α-anomeric nucleic acid molecule forms specific double-stranded hybrids with complementary RNA in which, contrary to the usual α-units, the strands run parallel to each other (Gaultier et al. (1987) Nucleic Acids Res 15: 6625-6641). The antisense nucleic acid molecule can also comprise a 2'-o-methylribonucleotide (Inoue et al. (1987) Nucleic Acids Res 15: 6131-6148) or a chimeric RNA -DNA analogue (Inoue et al. (1987) FEBS Lett 215: 327-330).

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4.4 RIBOZYMES AND PNA MOIETIES

In still another embodiment, an antisense nucleic acid of the invention is a ribozyme. Ribozymes are catalytic RNA molecules with ribonuclease activity that are capable of cleaving a single-stranded nucleic acid, such as an mRNA, to which they have a complementary region. Thus, ribozymes (e.g., hammerhead ribozymes (described in Haselhoff and Gerlach (1988) Nature 334:585-591)) can be used to catalytically cleave mRNA transcripts to thereby inhibit translation of an mRNA. A ribozyme having specificity for a nucleic acid of the invention can be designed based upon the nucleotide sequence of a DNA disclosed herein (i.e., SEQ ID NO: 1 - 438). For example, a derivative of Tetrahymena L-19 IVS RNA can be constructed in which the nucleotide sequence of the active site is complementary to the nucleotide sequence to be cleaved in a SECX-encoding mRNA. See, e.g., Cech et al. U.S. Pat. No. 4,987,071; and Cech et al. U.S. Pat. No. 5,116,742. Alternatively, SECX mRNA can be used to select a catalytic RNA having a specific ribonuclease activity from a pool of RNA molecules. See, e.g., Bartel et al., (1993) Science 261:1411-1418.

Alternatively, gene expression can be inhibited by targeting nucleotide sequences complementary to the regulatory region (e.g., promoter and/or enhancers) to form triple helical structures that prevent transcription of the gene in target cells. See generally, Helene. (1991) Anticancer Drug Des. 6: 569-84; Helene. et al. (1992) Ann. N.Y. Acad. Sci. 660:27-36; and Maher (1992) Bioassays 14: 807-15.

In various embodiments, the nucleic acids of the invention can be modified at the base moiety, sugar moiety or phosphate backbone to improve, e.g., the stability, hybridization, or solubility of the molecule. For example, the deoxyribose phosphate backbone of the nucleic acids can be modified to generate peptide nucleic acids (see

5 Hyrup et al. (1996) Bioorg Med Chem 4: 5-23). As used herein, the terms "peptide nucleic acids" or "PNAs" refer to nucleic acid mimics, e.g., DNA mimics, in which the deoxyribose phosphate backbone is replaced by a pseudopeptide backbone and only the four natural nucleobases are retained. The neutral backbone of PNAs has been shown to allow for specific hybridization to DNA and RNA under conditions of low ionic strength.

10 The synthesis of PNA oligomers can be performed using standard solid phase peptide synthesis protocols as described in Hyrup et al. (1996) above; Perry-O'Keefe et al. (1996) PNAS 93: 14670-675.

PNAs of the invention can be used in therapeutic and diagnostic applications. For example, PNAs can be used as antisense or antigene agents for sequence-specific modulation of gene expression by, e.g., inducing transcription or translation arrest or inhibiting replication. PNAs of the invention can also be used, e.g., in the analysis of single base pair mutations in a gene by, e.g., PNA directed PCR clamping; as artificial restriction enzymes when used in combination with other enzymes, e.g., S1 nucleases (Hyrup B. (1996) above); or as probes or primers for DNA sequence and hybridization (Hyrup et al. (1996), above; Perry-O'Keefe (1996), above).

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In another embodiment, PNAs of the invention can be modified, e.g., to enhance their stability or cellular uptake, by attaching lipophilic or other helper groups to PNA, by the formation of PNA-DNA chimeras, or by the use of liposomes or other techniques of drug delivery known in the art. For example, PNA-DNA chimeras can be generated that may combine the advantageous properties of PNA and DNA. Such chimeras allow DNA recognition enzymes, e.g., RNase H and DNA polymerases, to interact with the DNA portion while the PNA portion would provide high binding affinity and specificity. PNA-DNA chimeras can be linked using linkers of appropriate lengths selected in terms of base stacking, number of bonds between the nucleobases, and orientation (Hyrup (1996) above). The synthesis of PNA-DNA chimeras can be performed as described in Hyrup (1996) above and Finn et al. (1996) Nucl Acids Res 24: 3357-63. For example, a

DNA chain can be synthesized on a solid support using standard phosphoramidite coupling chemistry, and modified nucleoside analogs, e.g., 5'-(4-methoxytrityl)amino-5'-deoxy-thymidine phosphoramidite, can be used between the PNA and the 5' end of DNA (Mag et al. (1989) Nucl Acid Res 17: 5973-88). PNA monomers are then coupled in a stepwise manner to produce a chimeric molecule with a 5' PNA segment and a 3' DNA segment (Finn et al. (1996) above). Alternatively, chimeric molecules can be synthesized with a 5' DNA segment and a 3' PNA segment.

See, Petersen et al. (1975) Bioorg Med Chem Lett 5: 1119-11124.

In other embodiments, the oligonucleotide may include other appended groups such as peptides (e.g., for targeting host cell receptors in vivo), or agents facilitating transport across the cell membrane (see, e.g., Letsinger et al., 1989, Proc. Natl. Acad. Sci. U.S.A. 86:6553-6556; Lemaitre et al., 1987, Proc. Natl. Acad. Sci. 84:648-652; PCT Publication No. W088/09810) or the blood-brain barrier (see, e.g., PCT Publication No. W089/10134). In addition, oligonucleotides can be modified with hybridization triggered cleavage agents (See, e.g., Krol et al., 1988, BioTechniques 6:958-976) or intercalating agents. (See, e.g., Zon, 1988, Pharm. Res. 5: 539-549). To this end, the oligonucleotide may be conjugated to another molecule, e.g., a peptide, a hybridization triggered cross-linking agent, a transport agent, a hybridization-triggered cleavage agent, etc.

20 **4.5 HOSTS**

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The present invention further provides host cells genetically engineered to contain the polynucleotides of the invention. For example, such host cells may contain nucleic acids of the invention introduced into the host cell using known transformation, transfection or infection methods. The present invention still further provides host cells genetically engineered to express the polynucleotides of the invention, wherein such polynucleotides are in operative association with a regulatory sequence heterologous to the host cell which drives expression of the polynucleotides in the cell.

Knowledge of nucleic acid sequences allows for modification of cells to permit, or increase, expression of endogenous polypeptide. Cells can be modified (e.g., by homologous recombination) to provide increased polypeptide expression by replacing, in whole or in part, the naturally occurring promoter with all or part of a heterologous

promoter so that the cells express the polypeptide at higher levels. The heterologous promoter is inserted in such a manner that it is operatively linked to the encoding sequences. See, for example, PCT International Publication No. WO94/12650, PCT International Publication No. WO92/20808, and PCT International Publication No. WO91/09955. It is also contemplated that, in addition to heterologous promoter DNA, amplifiable marker DNA (e.g., ada, dhfr, and the multifunctional CAD gene which encodes carbamyl phosphate synthase, aspartate transcarbamylase, and dihydroorotase) and/or intron DNA may be inserted along with the heterologous promoter DNA. If linked to the coding sequence, amplification of the marker DNA by standard selection methods results in co-amplification of the desired protein coding sequences in the cells.

The host cell can be a higher eukaryotic host cell, such as a mammalian cell, a lower eukaryotic host cell, such as a yeast cell, or the host cell can be a prokaryotic cell, such as a bacterial cell. Introduction of the recombinant construct into the host cell can be effected by calcium phosphate transfection, DEAE, dextran mediated transfection, or electroporation (Davis, L. et al., *Basic Methods in Molecular Biology* (1986)). The host cells containing one of the polynucleotides of the invention, can be used in conventional manners to produce the gene product encoded by the isolated fragment (in the case of an ORF) or can be used to produce a heterologous protein under the control of the EMF.

Any host/vector system can be used to express one or more of the ORFs of the present invention. These include, but are not limited to, eukaryotic hosts such as HeLa cells, Cv-1 cell, COS cells, 293 cells, and Sf9 cells, as well as prokaryotic host such as E. coli and B. subtilis. The most preferred cells are those which do not normally express the particular polypeptide or protein or which expresses the polypeptide or protein at low natural level. Mature proteins can be expressed in mammalian cells, yeast, bacteria, or other cells under the control of appropriate promoters. Cell-free translation systems can also be employed to produce such proteins using RNAs derived from the DNA constructs of the present invention. Appropriate cloning and expression vectors for use with prokaryotic and eukaryotic hosts are described by Sambrook, et al., in Molecular Cloning: A Laboratory Manual, Second Edition, Cold Spring Harbor, New York (1989), the disclosure of which is hereby incorporated by reference.

Various mammalian cell culture systems can also be employed to express recombinant protein. Examples of mammalian expression systems include the COS-7 lines of monkey kidney fibroblasts, described by Gluzman, Cell 23:175 (1981). Other cell lines capable of expressing a compatible vector are, for example, the C127, monkey COS cells, Chinese Hamster Ovary (CHO) cells, human kidney 293 cells, human epidermal A431 cells, human Colo205 cells, 3T3 cells, CV-1 cells, other transformed primate cell lines, normal diploid cells, cell strains derived from in vitro culture of primary tissue, primary explants, HeLa cells, mouse L cells, BHK, HL-60, U937, HaK or Jurkat cells. Mammalian expression vectors will comprise an origin of replication, a suitable promoter and also any necessary ribosome binding sites, polyadenylation site, splice donor and acceptor sites, transcriptional termination sequences, and 5' flanking nontranscribed sequences. DNA sequences derived from the SV40 viral genome, for example, SV40 origin, early promoter, enhancer, splice, and polyadenylation sites may be used to provide the required nontranscribed genetic elements. Recombinant polypeptides and proteins produced in bacterial culture are usually isolated by initial extraction from cell pellets, followed by one or more salting-out, aqueous ion exchange or size exclusion chromatography steps. Protein refolding steps can be used, as necessary, in completing configuration of the mature protein. Finally, high performance liquid chromatography (HPLC) can be employed for final purification steps. Microbial cells employed in expression of proteins can be disrupted by any convenient method, including freeze-thaw cycling, sonication, mechanical disruption, or use of cell lysing agents.

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Alternatively, it may be possible to produce the protein in lower eukaryotes such as yeast or insects or in prokaryotes such as bacteria. Potentially suitable yeast strains include Saccharomyces cerevisiae, Schizosaccharomyces pombe, Kluyveromyces strains, Candida, or any yeast strain capable of expressing heterologous proteins. Potentially suitable bacterial strains include Escherichia coli, Bacillus subtilis, Salmonella typhimurium, or any bacterial strain capable of expressing heterologous proteins. If the protein is made in yeast or bacteria, it may be necessary to modify the protein produced therein, for example by phosphorylation or glycosylation of the appropriate sites, in order to obtain the functional protein. Such covalent attachments may be accomplished using known chemical or enzymatic methods.

In another embodiment of the present invention, cells and tissues may be engineered to express an endogenous gene comprising the polynucleotides of the invention under the control of inducible regulatory elements, in which case the regulatory sequences of the endogenous gene may be replaced by homologous recombination. As described herein, gene targeting can be used to replace a gene's existing regulatory region with a regulatory sequence isolated from a different gene or a novel regulatory sequence synthesized by genetic engineering methods. Such regulatory sequences may be comprised of promoters, enhancers, scaffold-attachment regions, negative regulatory elements, transcriptional initiation sites, regulatory protein binding sites or combinations of said sequences. Alternatively, sequences which affect the structure or stability of the RNA or protein produced may be replaced, removed, added, or otherwise modified by targeting. These sequence include polyadenylation signals, mRNA stability elements, splice sites, leader sequences for enhancing or modifying transport or secretion properties of the protein, or other sequences which alter or improve the function or stability of protein or RNA molecules.

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The targeting event may be a simple insertion of the regulatory sequence, placing the gene under the control of the new regulatory sequence, e.g., inserting a new promoter or enhancer or both upstream of a gene. Alternatively, the targeting event may be a simple deletion of a regulatory element, such as the deletion of a tissue-specific negative regulatory element. Alternatively, the targeting event may replace an existing element; for example, a tissue-specific enhancer can be replaced by an enhancer that has broader or different cell-type specificity than the naturally occurring elements. Here, the naturally occurring sequences are deleted and new sequences are added. In all cases, the identification of the targeting event may be facilitated by the use of one or more selectable marker genes that are contiguous with the targeting DNA, allowing for the selection of cells in which the exogenous DNA has integrated into the host cell genome. The identification of the targeting event may also be facilitated by the use of one or more marker genes exhibiting the property of negative selection, such that the negatively selectable marker is linked to the exogenous DNA, but configured such that the negatively selectable marker flanks the targeting sequence, and such that a correct homologous recombination event with sequences in the host cell genome does not result

in the stable integration of the negatively selectable marker. Markers useful for this purpose include the Herpes Simplex Virus thymidine kinase (TK) gene or the bacterial xanthine-guanine phosphoribosyl-transferase (gpt) gene.

The gene targeting or gene activation techniques which can be used in accordance with this aspect of the invention are more particularly described in U.S. Patent No. 5,272,071 to Chappel; U.S. Patent No. 5,578,461 to Sherwin et al.; International Application No. PCT/US92/09627 (WO93/09222) by Selden et al.; and International Application No. PCT/US90/06436 (WO91/06667) by Skoultchi et al., each of which is incorporated by reference herein in its entirety.

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4.6 POLYPEPTIDES OF THE INVENTION

The isolated polypeptides of the invention include, but are not limited to, a polypeptide comprising: the amino acid sequences set forth as any one of SEQ ID NO: 1-438 or an amino acid sequence encoded by any one of the nucleotide sequences SEQ ID NOs: 1 - 438 or the corresponding full length or mature protein. Polypeptides of the invention also include polypeptides preferably with biological or immunological activity that are encoded by: (a) a polynucleotide having any one of the nucleotide sequences set forth in SEQ ID NOs: 1 – 438 or (b) polynucleotides encoding any one of the amino acid sequences set forth as SEQ ID NO: 1-438 or (c) polynucleotides that hybridize to the complement of the polynucleotides of either (a) or (b) under stringent hybridization conditions. The invention also provides biologically active or immunologically active variants of any of the amino acid sequences set forth as SEQ ID NO: 1-438 or the corresponding full length or mature protein; and "substantial equivalents" thereof (e.g., with at least about 65%, at least about 70%, at least about 75%, at least about 80%, at least about 85%, 86%, 87%, 88%, 89%, at least about 90%, 91%, 92%, 93%, 94%, typically at least about 95%, 96%, 97%, more typically at least about 98%, or most typically at least about 99% amino acid identity) that retain biological activity. Polypeptides encoded by allelic variants may have a similar, increased, or decreased activity compared to polypeptides comprising SEQ ID NO: 1-438.

Fragments of the proteins of the present invention which are capable of exhibiting biological activity are also encompassed by the present invention. Fragments of the

protein may be in linear form or they may be cyclized using known methods, for example, as described in H. U. Saragovi, et al., Bio/Technology 10, 773-778 (1992) and in R. S. McDowell, et al., J. Amer. Chem. Soc. 114, 9245-9253 (1992), both of which are incorporated herein by reference. Such fragments may be fused to carrier molecules such as immunoglobulins for many purposes, including increasing the valency of protein binding sites.

The present invention also provides both full-length and mature forms (for example, without a signal sequence or precursor sequence) of the disclosed proteins. The protein coding sequence is identified in the sequence listing by translation of the disclosed nucleotide sequences. The mature form of such protein may be obtained by expression of a full-length polynucleotide in a suitable mammalian cell or other host cell. The sequence of the mature form of the protein is also determinable from the amino acid sequence of the full-length form. Where proteins of the present invention are membrane bound, soluble forms of the proteins are also provided. In such forms, part or all of the regions causing the proteins to be membrane bound are deleted so that the proteins are fully secreted from the cell in which they are expressed.

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Protein compositions of the present invention may further comprise an acceptable carrier, such as a hydrophilic, e.g., pharmaceutically acceptable, carrier.

The present invention further provides isolated polypeptides encoded by the nucleic acid fragments of the present invention or by degenerate variants of the nucleic acid fragments of the present invention. By "degenerate variant" is intended nucleotide fragments which differ from a nucleic acid fragment of the present invention (e.g., an ORF) by nucleotide sequence but, due to the degeneracy of the genetic code, encode an identical polypeptide sequence. Preferred nucleic acid fragments of the present invention are the ORFs that encode proteins.

A variety of methodologies known in the art can be utilized to obtain any one of the isolated polypeptides or proteins of the present invention. At the simplest level, the amino acid sequence can be synthesized using commercially available peptide synthesizers. The synthetically-constructed protein sequences, by virtue of sharing primary, secondary or tertiary structural and/or conformational characteristics with proteins may possess biological properties in common therewith, including protein

activity. This technique is particularly useful in producing small peptides and fragments of larger polypeptides. Fragments are useful, for example, in generating antibodies against the native polypeptide. Thus, they may be employed as biologically active or immunological substitutes for natural, purified proteins in screening of therapeutic compounds and in immunological processes for the development of antibodies.

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The polypeptides and proteins of the present invention can alternatively be purified from cells which have been altered to express the desired polypeptide or protein. As used herein, a cell is said to be altered to express a desired polypeptide or protein when the cell, through genetic manipulation, is made to produce a polypeptide or protein which it normally does not produce or which the cell normally produces at a lower level. One skilled in the art can readily adapt procedures for introducing and expressing either recombinant or synthetic sequences into eukaryotic or prokaryotic cells in order to generate a cell which produces one of the polypeptides or proteins of the present invention.

The invention also relates to methods for producing a polypeptide comprising growing a culture of host cells of the invention in a suitable culture medium, and purifying the protein from the cells or the culture in which the cells are grown. For example, the methods of the invention include a process for producing a polypeptide in which a host cell containing a suitable expression vector that includes a polynucleotide of the invention is cultured under conditions that allow expression of the encoded polypeptide. The polypeptide can be recovered from the culture, conveniently from the culture medium, or from a lysate prepared from the host cells and further purified. Preferred embodiments include those in which the protein produced by such process is a full length or mature form of the protein.

In an alternative method, the polypeptide or protein is purified from bacterial cells which naturally produce the polypeptide or protein. One skilled in the art can readily follow known methods for isolating polypeptides and proteins in order to obtain one of the isolated polypeptides or proteins of the present invention. These include, but are not limited to, immunochromatography, HPLC, size-exclusion chromatography, ion-exchange chromatography, and immuno-affinity chromatography. See, e.g., Scopes, Protein Purification: Principles and Practice, Springer-Verlag (1994); Sambrook, et al.,

in Molecular Cloning: A Laboratory Manual; Ausubel et al., Current Protocols in Molecular Biology. Polypeptide fragments that retain biological/immunological activity include fragments comprising greater than about 100 amino acids, or greater than about 200 amino acids, and fragments that encode specific protein domains.

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The purified polypeptides can be used in *in vitro* binding assays which are well known in the art to identify molecules which bind to the polypeptides. These molecules include but are not limited to, for e.g., small molecules, molecules from combinatorial libraries, antibodies or other proteins. The molecules identified in the binding assay are then tested for antagonist or agonist activity in *in vivo* tissue culture or animal models that are well known in the art. In brief, the molecules are titrated into a plurality of cell cultures or animals and then tested for either cell/animal death or prolonged survival of the animal/cells.

In addition, the peptides of the invention or molecules capable of binding to the peptides may be complexed with toxins, e.g., ricin or cholera, or with other compounds that are toxic to cells. The toxin-binding molecule complex is then targeted to a tumor or other cell by the specificity of the binding molecule for SEQ ID NO: 1-438.

The protein of the invention may also be expressed as a product of transgenic animals, e.g., as a component of the milk of transgenic cows, goats, pigs, or sheep which are characterized by somatic or germ cells containing a nucleotide sequence encoding the protein.

The proteins provided herein also include proteins characterized by amino acid sequences similar to those of purified proteins but into which modification are naturally provided or deliberately engineered. For example, modifications, in the peptide or DNA sequence, can be made by those skilled in the art using known techniques. Modifications of interest in the protein sequences may include the alteration, substitution, replacement, insertion or deletion of a selected amino acid residue in the coding sequence. For example, one or more of the cysteine residues may be deleted or replaced with another amino acid to alter the conformation of the molecule. Techniques for such alteration, substitution, replacement, insertion or deletion are well known to those skilled in the art (see, e.g., U.S. Pat. No. 4,518,584). Preferably, such alteration, substitution, replacement, insertion or deletion retains the desired activity of the protein. Regions of the protein that

are important for the protein function can be determined by various methods known in the art including the alanine-scanning method which involved systematic substitution of single or strings of amino acids with alanine, followed by testing the resulting alanine-containing variant for biological activity. This type of analysis determines the importance of the substituted amino acid(s) in biological activity. Regions of the protein that are important for protein function may be determined by the eMATRIX program.

Other fragments and derivatives of the sequences of proteins which would be expected to retain protein activity in whole or in part and are useful for screening or other immunological methodologies may also be easily made by those skilled in the art given the disclosures herein. Such modifications are encompassed by the present invention.

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The protein may also be produced by operably linking the isolated polynucleotide of the invention to suitable control sequences in one or more insect expression vectors, and employing an insect expression system. Materials and methods for baculovirus/insect cell expression systems are commercially available in kit form from, e.g., Invitrogen, San Diego, Calif., U.S.A. (the MaxBatTM kit), and such methods are well known in the art, as described in Summers and Smith, Texas Agricultural Experiment Station Bulletin No. 1555 (1987), incorporated herein by reference. As used herein, an insect cell capable of expressing a polynucleotide of the present invention is "transformed."

The protein of the invention may be prepared by culturing transformed host cells under culture conditions suitable to express the recombinant protein. The resulting expressed protein may then be purified from such culture (*i.e.*, from culture medium or cell extracts) using known purification processes, such as gel filtration and ion exchange chromatography. The purification of the protein may also include an affinity column containing agents which will bind to the protein; one or more column steps over such affinity resins as concanavalin A-agarose, heparin-toyopearlTM or Cibacrom blue 3GA SepharoseTM; one or more steps involving hydrophobic interaction chromatography using such resins as phenyl ether, butyl ether, or propyl ether; or immunoaffinity chromatography.

Alternatively, the protein of the invention may also be expressed in a form which will facilitate purification. For example, it may be expressed as a fusion protein, such as

those of maltose binding protein (MBP), glutathione-S-transferase (GST) or thioredoxin (TRX), or as a His tag. Kits for expression and purification of such fusion proteins are commercially available from New England BioLab (Beverly, Mass.), Pharmacia (Piscataway, N.J.) and Invitrogen, respectively. The protein can also be tagged with an epitope and subsequently purified by using a specific antibody directed to such epitope. One such epitope ("FLAG®") is commercially available from Kodak (New Haven, Conn.).

Finally, one or more reverse-phase high performance liquid chromatography (RP-HPLC) steps employing hydrophobic RP-HPLC media, e.g., silica gel having pendant methyl or other aliphatic groups, can be employed to further purify the protein. Some or all of the foregoing purification steps, in various combinations, can also be employed to provide a substantially homogeneous isolated recombinant protein. The protein thus purified is substantially free of other mammalian proteins and is defined in accordance with the present invention as an "isolated protein."

The polypeptides of the invention include analogs (variants). This embraces fragments, as well as peptides in which one or more amino acids has been deleted, inserted, or substituted. Also, analogs of the polypeptides of the invention embrace fusions of the polypeptides or modifications of the polypeptides of the invention, wherein the polypeptide or analog is fused to another moiety or moieties, e.g., targeting moiety or another therapeutic agent. Such analogs may exhibit improved properties such as activity and/or stability. Examples of moieties which may be fused to the polypeptide or an analog include, for example, targeting moieties which provide for the delivery of polypeptide to pancreatic cells, e.g., antibodies to pancreatic cells, antibodies to immune cells such as T-cells, monocytes, dendritic cells, granulocytes, etc., as well as receptor and ligands expressed on pancreatic or immune cells. Other moieties which may be fused to the polypeptide include therapeutic agents which are used for treatment, for example, immunosuppressive drugs such as cyclosporin, SK506, azathioprine, CD3 antibodies and steroids. Also, polypeptides may be fused to immune modulators, and other cytokines such as alpha or beta interferon.

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4.6.1 DETERMINING POLYPEPTIDE AND POLYNUCLEOTIDE IDENTITY AND SIMILARITY

Preferred identity and/or similarity are designed to give the largest match between the sequences tested. Methods to determine identity and similarity are codified in computer programs including, but are not limited to, the GCG program package, including GAP (Devereux, J., et al., Nucleic Acids Research 12(1):387 (1984); Genetics Computer Group, University of Wisconsin, Madison, WI), BLASTP, BLASTN, BLASTX, FASTA (Altschul, S.F. et al., J. Molec. Biol. 215:403-410 (1990), PSI-BLAST (Altschul S.F. et al., Nucleic Acids Res. vol. 25, pp. 3389-3402, herein incorporated by 10 reference), eMatrix software (Wu et al., J. Comp. Biol., Vol. 6, pp. 219-235 (1999), herein incorporated by reference), eMotif software (Nevill-Manning et al, ISMB-97, Vol. 4, pp. 202-209, herein incorporated by reference), pFam software (Sonnhammer et al., Nucleic Acids Res., Vol. 26(1), pp. 320-322 (1998), herein incorporated by reference) and the Kyte-Doolittle hydrophobocity prediction algorithm (J. Mol Biol, 157, pp. 105-31 (1982), incorporated herein by reference). The BLAST programs are publicly available 15 from the National Center for Biotechnology Information (NCBI) and other sources (BLAST Manual, Altschul, S., et al. NCB NLM NIH Bethesda, MD 20894; Altschul, S., et al., J. Mol. Biol. 215:403-410 (1990).

4.7 CHIMERIC AND FUSION PROTEINS

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The invention also provides chimeric or fusion proteins. As used herein, a "chimeric protein" or "fusion protein" comprises a polypeptide of the invention operatively linked to another polypeptide. Within a fusion protein the polypeptide according to the invention can correspond to all or a portion of a protein according to the invention. In one embodiment, a fusion protein comprises at least one biologically active portion of a protein according to the invention. In another embodiment, a fusion protein comprises at least two biologically active portions of a protein according to the invention. Within the fusion protein, the term "operatively linked" is intended to indicate that the polypeptide according to the invention and the other polypeptide are fused in-frame to each other. The polypeptide can be fused to the N-terminus or C-terminus, or to the middle.

For example, in one embodiment a fusion protein comprises a polypeptide according to the invention operably linked to the extracellular domain of a second protein.

In another embodiment, the fusion protein is a GST-fusion protein in which the polypeptide sequences of the invention are fused to the C-terminus of the GST (i.e., glutathione S-transferase) sequences.

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In another embodiment, the fusion protein is an immunoglobulin fusion protein in which the polypeptide sequences according to the invention comprise one or more domains fused to sequences derived from a member of the immunoglobulin protein family. The immunoglobulin fusion proteins of the invention can be incorporated into pharmaceutical compositions and administered to a subject to inhibit an interaction between a ligand and a protein of the invention on the surface of a cell, to thereby suppress signal transduction *in vivo*. The immunoglobulin fusion proteins can be used to affect the bioavailability of a cognate ligand. Inhibition of the ligand/protein interaction may be useful therapeutically for both the treatment of proliferative and differentiative disorders, *e.g.*, cancer as well as modulating (*e.g.*, promoting or inhibiting) cell survival. Moreover, the immunoglobulin fusion proteins of the invention can be used as immunogens to produce antibodies in a subject, to purify ligands, and in screening assays to identify molecules that inhibit the interaction of a polypeptide of the invention with a ligand.

A chimeric or fusion protein of the invention can be produced by standard recombinant DNA techniques. For example, DNA fragments coding for the different polypeptide sequences are ligated together in-frame in accordance with conventional techniques, e.g., by employing blunt-ended or stagger-ended termini for ligation, restriction enzyme digestion to provide for appropriate termini, filling-in of cohesive ends as appropriate, alkaline phosphatase treatment to avoid undesirable joining, and enzymatic ligation. In another embodiment, the fusion gene can be synthesized by conventional techniques including automated DNA synthesizers. Alternatively, PCR amplification of gene fragments can be carried out using anchor primers that give rise to complementary overhangs between two consecutive gene fragments that can subsequently be annealed and reamplified to generate a chimeric gene sequence (see, for

example, Ausubel et al. (eds.) CURRENT PROTOCOLS IN MOLECULAR BIOLOGY, John Wiley & Sons, 1992). Moreover, many expression vectors are commercially available that already encode a fusion moiety (e.g., a GST polypeptide). A nucleic acid encoding a polypeptide of the invention can be cloned into such an expression vector such that the fusion moiety is linked in-frame to the protein of the invention.

4.8 GENE THERAPY

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Mutations in the polynucleotides of the invention gene may result in loss of normal function of the encoded protein. The invention thus provides gene therapy to 10 restore normal activity of the polypeptides of the invention; or to treat disease states involving polypeptides of the invention. Delivery of a functional gene encoding polypeptides of the invention to appropriate cells is effected ex vivo, in situ, or in vivo by use of vectors, and more particularly viral vectors (e.g., adenovirus, adeno-associated virus, or a retrovirus), or ex vivo by use of physical DNA transfer methods (e.g., liposomes or chemical treatments). See, for example, Anderson, Nature, supplement to 15 vol. 392, no. 6679, pp.25-20 (1998). For additional reviews of gene therapy technology see Friedmann, Science, 244: 1275-1281 (1989); Verma, Scientific American: 68-84 (1990); and Miller, Nature, 357: 455-460 (1992). Introduction of any one of the nucleotides of the present invention or a gene encoding the polypeptides of the present invention can also be accomplished with extrachromosomal substrates (transient expression) or artificial chromosomes (stable expression). Cells may also be cultured ex vivo in the presence of proteins of the present invention in order to proliferate or to produce a desired effect on or activity in such cells. Treated cells can then be introduced in vivo for therapeutic purposes. Alternatively, it is contemplated that in other human disease states, preventing the expression of or inhibiting the activity of polypeptides of the invention will be useful in treating the disease states. It is contemplated that antisense therapy or gene therapy could be applied to negatively regulate the expression of polypeptides of the invention.

Other methods inhibiting expression of a protein include the introduction of antisense molecules to the nucleic acids of the present invention, their complements, or their translated RNA sequences, by methods known in the art. Further, the polypeptides of the

present invention can be inhibited by using targeted deletion methods, or the insertion of a negative regulatory element such as a silencer, which is tissue specific.

The present invention still further provides cells genetically engineered *in vivo* to express the polynucleotides of the invention, wherein such polynucleotides are in operative association with a regulatory sequence heterologous to the host cell which drives expression of the polynucleotides in the cell. These methods can be used to increase or decrease the expression of the polynucleotides of the present invention.

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Knowledge of DNA sequences provided by the invention allows for modification of cells to permit, increase, or decrease, expression of endogenous polypeptide. Cells can be modified (e.g., by homologous recombination) to provide increased polypeptide expression by replacing, in whole or in part, the naturally occurring promoter with all or part of a heterologous promoter so that the cells express the protein at higher levels. The heterologous promoter is inserted in such a manner that it is operatively linked to the desired protein encoding sequences. See, for example, PCT International Publication No. WO 94/12650, PCT International Publication No. WO 92/20808, and PCT International Publication No. WO 91/09955. It is also contemplated that, in addition to heterologous promoter DNA, amplifiable marker DNA (e.g., ada, dhfr, and the multifunctional CAD gene which encodes carbamyl phosphate synthase, aspartate transcarbamylase, and dihydroorotase) and/or intron DNA may be inserted along with the heterologous promoter DNA. If linked to the desired protein coding sequence, amplification of the marker DNA by standard selection methods results in co-amplification of the desired protein coding sequences in the cells.

In another embodiment of the present invention, cells and tissues may be engineered to express an endogenous gene comprising the polynucleotides of the invention under the control of inducible regulatory elements, in which case the regulatory sequences of the endogenous gene may be replaced by homologous recombination. As described herein, gene targeting can be used to replace a gene's existing regulatory region with a regulatory sequence isolated from a different gene or a novel regulatory sequence synthesized by genetic engineering methods. Such regulatory sequences may be comprised of promoters, enhancers, scaffold-attachment regions, negative regulatory elements, transcriptional initiation sites, regulatory protein binding sites or combinations of said sequences.

Alternatively, sequences which affect the structure or stability of the RNA or protein

produced may be replaced, removed, added, or otherwise modified by targeting. These sequences include polyadenylation signals, mRNA stability elements, splice sites, leader sequences for enhancing or modifying transport or secretion properties of the protein, or other sequences which alter or improve the function or stability of protein or RNA molecules.

The targeting event may be a simple insertion of the regulatory sequence, placing the gene under the control of the new regulatory sequence, e.g., inserting a new promoter or enhancer or both upstream of a gene. Alternatively, the targeting event may be a simple deletion of a regulatory element, such as the deletion of a tissue-specific negative regulatory element. Alternatively, the targeting event may replace an existing element; for example, a 10 tissue-specific enhancer can be replaced by an enhancer that has broader or different cell-type specificity than the naturally occurring elements. Here, the naturally occurring sequences are deleted and new sequences are added. In all cases, the identification of the targeting event may be facilitated by the use of one or more selectable marker genes that are 15 contiguous with the targeting DNA, allowing for the selection of cells in which the exogenous DNA has integrated into the cell genome. The identification of the targeting event may also be facilitated by the use of one or more marker genes exhibiting the property of negative selection, such that the negatively selectable marker is linked to the exogenous DNA, but configured such that the negatively selectable marker flanks the targeting sequence, and such that a correct homologous recombination event with sequences in the host cell genome does not result in the stable integration of the negatively selectable marker. Markers useful for this purpose include the Herpes Simplex Virus thymidine kinase (TK) gene or the bacterial xanthine-guanine phosphoribosyl-transferase (gpt) gene.

The gene targeting or gene activation techniques which can be used in accordance with this aspect of the invention are more particularly described in U.S. Patent No. 5,272,071 to Chappel; U.S. Patent No. 5,578,461 to Sherwin et al.; International Application No. PCT/US92/09627 (WO93/09222) by Selden et al.; and International Application No. PCT/US90/06436 (WO91/06667) by Skoultchi et al., each of which is incorporated by reference herein in its entirety.

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4.9 TRANSGENIC ANIMALS

In preferred methods to determine biological functions of the polypeptides of theinvention in vivo, one or more genes provided by the invention are either over expressed
or inactivated in the germ line of animals using homologous recombination [Capecchi,
Science 244:1288-1292 (1989)]. Animals in which the gene is over expressed, under the
regulatory control of exogenous or endogenous promoter elements, are known as
transgenic animals. Animals in which an endogenous gene has been inactivated by
homologous recombination are referred to as "knockout" animals. Knockout animals,
preferably non-human mammals, can be prepared as described in U.S. Patent No.
5,557,032, incorporated herein by reference. Transgenic animals are useful to determine
the roles polypeptides of the invention play in biological processes, and preferably in
disease states. Transgenic animals are useful as model systems to identify compounds
that modulate lipid metabolism. Transgenic animals, preferably non-human mammals,
are produced using methods as described in U.S. Patent No 5,489,743 and PCT
Publication No. WO94/28122, incorporated herein by reference.

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Transgenic animals can be prepared wherein all or part of a promoter of the polynucleotides of the invention is either activated or inactivated to alter the level of expression of the polypeptides of the invention. Inactivation can be carried out using homologous recombination methods described above. Activation can be achieved by supplementing or even replacing the homologous promoter to provide for increased protein expression. The homologous promoter can be supplemented by insertion of one or more heterologous enhancer elements known to confer promoter activation in a particular tissue.

The polynucleotides of the present invention also make possible the development, through, e.g., homologous recombination or knock out strategies, of animals that fail to express polypeptides of the invention or that express a variant polypeptide. Such animals are useful as models for studying the *in vivo* activities of polypeptide as well as for studying modulators of the polypeptides of the invention.

In preferred methods to determine biological functions of the polypeptides of the invention *in vivo*, one or more genes provided by the invention are either over expressed or inactivated in the germ line of animals using homologous recombination [Capecchi, Science 244:1288-1292 (1989)]. Animals in which the gene is over expressed, under the

regulatory control of exogenous or endogenous promoter elements, are known as transgenic animals. Animals in which an endogenous gene has been inactivated by homologous recombination are referred to as "knockout" animals. Knockout animals, preferably non-human mammals, can be prepared as described in U.S. Patent No. 5,557,032, incorporated herein by reference. Transgenic animals are useful to determine the roles polypeptides of the invention play in biological processes, and preferably in disease states. Transgenic animals are useful as model systems to identify compounds that modulate lipid metabolism. Transgenic animals, preferably non-human mammals, are produced using methods as described in U.S. Patent No 5,489,743 and PCT Publication No. WO94/28122, incorporated herein by reference.

Transgenic animals can be prepared wherein all or part of the polynucleotides of the invention promoter is either activated or inactivated to alter the level of expression of the polypeptides of the invention. Inactivation can be carried out using homologous recombination methods described above. Activation can be achieved by supplementing or even replacing the homologous promoter to provide for increased protein expression. The homologous promoter can be supplemented by insertion of one or more heterologous enhancer elements known to confer promoter activation in a particular tissue.

4.10 USES AND BIOLOGICAL ACTIVITY

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The polynucleotides and proteins of the present invention are expected to exhibit one or more of the uses or biological activities (including those associated with assays cited herein) identified herein. Uses or activities described for proteins of the present invention may be provided by administration or use of such proteins or of polynucleotides encoding such proteins (such as, for example, in gene therapies or vectors suitable for introduction of DNA). The mechanism underlying the particular condition or pathology will dictate whether the polypeptides of the invention, the polynucleotides of the invention or modulators (activators or inhibitors) thereof would be beneficial to the subject in need of treatment. Thus, "therapeutic compositions of the invention" include compositions comprising isolated polynucleotides (including recombinant DNA molecules, cloned genes and degenerate variants thereof) or polypeptides of the invention (including full length protein, mature protein and

truncations or domains thereof), or compounds and other substances that modulate the overall activity of the target gene products, either at the level of target gene/protein expression or target protein activity. Such modulators include polypeptides, analogs, (variants), including fragments and fusion proteins, antibodies and other binding proteins; chemical compounds that directly or indirectly activate or inhibit the polypeptides of the invention (identified, e.g., via drug screening assays as described herein); antisense polynucleotides and polynucleotides suitable for triple helix formation; and in particular antibodies or other binding partners that specifically recognize one or more epitopes of the polypeptides of the invention.

The polypeptides of the present invention may likewise be involved in cellular activation or in one of the other physiological pathways described herein.

4.10.1 RESEARCH USES AND UTILITIES

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The polynucleotides provided by the present invention can be used by the research community for various purposes. The polynucleotides can be used to express recombinant protein for analysis, characterization or therapeutic use; as markers for tissues in which the corresponding protein is preferentially expressed (either constitutively or at a particular stage of tissue differentiation or development or in disease states); as molecular weight markers on gels; as chromosome markers or tags (when labeled) to identify chromosomes or to map related gene positions; to compare with endogenous DNA sequences in patients to identify potential genetic disorders; as probes to hybridize and thus discover novel, related DNA sequences; as a source of information to derive PCR primers for genetic fingerprinting; as a probe to "subtract-out" known sequences in the process of discovering other novel polynucleotides; for selecting and making oligomers for attachment to a "gene chip" or other support, including for examination of expression patterns; to raise anti-protein antibodies using DNA immunization techniques; and as an antigen to raise anti-DNA antibodies or elicit another immune response. Where the polynucleotide encodes a protein which binds or potentially binds to another protein (such as, for example, in a receptor-ligand interaction), the polynucleotide can also be used in interaction trap assays (such as, for example, that described in Gyuris et al., Cell 75:791-803 (1993)) to identify

polynucleotides encoding the other protein with which binding occurs or to identify inhibitors of the binding interaction.

The polypeptides provided by the present invention can similarly be used in assays to determine biological activity, including in a panel of multiple proteins for high-throughput screening; to raise antibodies or to elicit another immune response; as a reagent (including the labeled reagent) in assays designed to quantitatively determine levels of the protein (or its receptor) in biological fluids; as markers for tissues in which the corresponding polypeptide is preferentially expressed (either constitutively or at a particular stage of tissue differentiation or development or in a disease state); and, of course, to isolate correlative receptors or ligands. Proteins involved in these binding interactions can also be used to screen for peptide or small molecule inhibitors or agonists of the binding interaction.

Any or all of these research utilities are capable of being developed into reagent grade or kit format for commercialization as research products.

Methods for performing the uses listed above are well known to those skilled in the art. References disclosing such methods include without limitation "Molecular Cloning: A Laboratory Manual", 2d ed., Cold Spring Harbor Laboratory Press, Sambrook, J., E. F. Fritsch and T. Maniatis eds., 1989, and "Methods in Enzymology: Guide to Molecular Cloning Techniques", Academic Press, Berger, S. L. and A. R.
Kimmel eds., 1987.

4.10.2 NUTRITIONAL USES

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Polynucleotides and polypeptides of the present invention can also be used as nutritional sources or supplements. Such uses include without limitation use as a protein or amino acid supplement, use as a carbon source, use as a nitrogen source and use as a source of carbohydrate. In such cases the polypeptide or polynucleotide of the invention can be added to the feed of a particular organism or can be administered as a separate solid or liquid preparation, such as in the form of powder, pills, solutions, suspensions or capsules. In the case of microorganisms, the polypeptide or polynucleotide of the invention can be added to the medium in or on which the microorganism is cultured.

4.10.3 CYTOKINE AND CELL PROLIFERATION/DIFFERENTIATION ACTIVITY

A polypeptide of the present invention may exhibit activity relating to cytokine, cell proliferation (either inducing or inhibiting) or cell differentiation (either inducing or inhibiting) activity or may induce production of other cytokines in certain cell populations. A polynucleotide of the invention can encode a polypeptide exhibiting such attributes. Many protein factors discovered to date, including all known cytokines, have exhibited activity in one or more factor-dependent cell proliferation assays, and hence the assays serve as a convenient confirmation of cytokine activity. The activity of therapeutic compositions of the present invention is evidenced by any one of a number of routine factor dependent cell proliferation assays for cell lines including, without limitation, 32D, DA2, DA1G, T10, B9, B9/11, BaF3, MC9/G, M+(preB M+), 2E8, RB5, DA1, 123, T1165, HT2, CTLL2, TF-1, Mo7e, CMK, HUVEC, and Caco. Therapeutic compositions of the invention can be used in the following:

Assays for T-cell or thymocyte proliferation include without limitation those described in: Current Protocols in Immunology, Ed by J. E. Coligan, A. M. Kruisbeek, D. H. Margulies, E. M. Shevach, W. Strober, Pub. Greene Publishing Associates and Wiley-Interscience (Chapter 3, *In Vitro* assays for Mouse Lymphocyte Function 3.1-3.19; Chapter 7, Immunologic studies in Humans); Takai et al., J. Immunol. 137:3494-3500,
1986; Bertagnolli et al., J. Immunol. 145:1706-1712, 1990; Bertagnolli et al., Cellular Immunology 133:327-341, 1991; Bertagnolli, et al., I. Immunol. 149:3778-3783, 1992; Bowman et al., I. Immunol. 152:1756-1761, 1994.

Assays for cytokine production and/or proliferation of spleen cells, lymph node cells or thymocytes include, without limitation, those described in: Polyclonal T cell stimulation, Kruisbeek, A. M. and Shevach, E. M. In Current Protocols in Immunology. J. E. e.a. Coligan eds. Vol 1 pp. 3.12.1-3.12.14, John Wiley and Sons, Toronto. 1994; and Measurement of mouse and human interleukin-γ, Schreiber, R. D. In Current Protocols in Immunology. J. E. e.a. Coligan eds. Vol 1 pp. 6.8.1-6.8.8, John Wiley and Sons, Toronto. 1994.

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Assays for proliferation and differentiation of hematopoietic and lymphopoietic cells include, without limitation, those described in: Measurement of Human and Murine

Interleukin 2 and Interleukin 4, Bottomly, K., Davis, L. S. and Lipsky, P. E. In Current Protocols in Immunology. J. E. e.a. Coligan eds. Vol 1 pp. 6.3.1-6.3.12, John Wiley and Sons, Toronto. 1991; deVries et al., J. Exp. Med. 173:1205-1211, 1991; Moreau et al., Nature 336:690-692, 1988; Greenberger et al., Proc. Natl. Acad. Sci. U.S.A.

- 80:2931-2938, 1983; Measurement of mouse and human interleukin 6--Nordan, R. In Current Protocols in Immunology. J. E. Coligan eds. Vol 1 pp. 6.6.1-6.6.5, John Wiley and Sons, Toronto. 1991; Smith et al., Proc. Natl. Aced. Sci. U.S.A. 83:1857-1861, 1986; Measurement of human Interleukin 11--Bennett, F., Giannotti, J., Clark, S. C. and Turner, K. J. In Current Protocols in Immunology. J. E. Coligan eds. Vol 1 pp. 6.15.1 John
- Wiley and Sons, Toronto. 1991; Measurement of mouse and human Interleukin 9--Ciarletta, A., Giannotti, J., Clark, S. C. and Turner, K. J. In Current Protocols in Immunology. J. E. Coligan eds. Vol 1 pp. 6.13.1, John Wiley and Sons, Toronto. 1991.

Assays for T-cell clone responses to antigens (which will identify, among others, proteins that affect APC-T cell interactions as well as direct T-cell effects by measuring proliferation and cytokine production) include, without limitation, those described in: Current Protocols in Immunology, Ed by J. E. Coligan, A. M. Kruisbeek, D. H. Margulies, E. M. Shevach, W Strober, Pub. Greene Publishing Associates and Wiley-Interscience (Chapter 3, *In Vitro* assays for Mouse Lymphocyte Function; Chapter 6, Cytokines and their cellular receptors; Chapter 7, Immunologic studies in Humans); Weinberger et al., Proc. Natl. Acad. Sci. USA 77:6091-6095, 1980; Weinberger et al.,

Weinberger et al., Proc. Natl. Acad. Sci. USA 77:6091-6095, 1980; Weinberger et al.,
 Eur. J. Immun. 11:405-411, 1981; Takai et al., J. Immunol. 137:3494-3500, 1986; Takai et al., J. Immunol. 140:508-512, 1988.

4.10.4 STEM CELL GROWTH FACTOR ACTIVITY

A polypeptide of the present invention may exhibit stem cell growth factor activity and be involved in the proliferation, differentiation and survival of pluripotent and totipotent stem cells including primordial germ cells, embryonic stem cells, hematopoietic stem cells and/or germ line stem cells. Administration of the polypeptide of the invention to stem cells in vivo or ex vivo is expected to maintain and expand cell populations in a totipotential or pluripotential state which would be useful for reengineering damaged or diseased tissues, transplantation, manufacture of bio-

pharmaceuticals and the development of bio-sensors. The ability to produce large quantities of human cells has important working applications for the production of human proteins which currently must be obtained from non-human sources or donors, implantation of cells to treat diseases such as Parkinson's, Alzheimer's and other neurodegenerative diseases; tissues for grafting such as bone marrow, skin, cartilage, tendons, bone, muscle (including cardiac muscle), blood vessels, cornea, neural cells, gastrointestinal cells and others; and organs for transplantation such as kidney, liver, pancreas (including islet cells), heart and lung.

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It is contemplated that multiple different exogenous growth factors and/or cytokines may be administered in combination with the polypeptide of the invention to achieve the desired effect, including any of the growth factors listed herein, other stem cell maintenance factors, and specifically including stem cell factor (SCF), leukemia inhibitory factor (LIF), Flt-3 ligand (Flt-3L), any of the interleukins, recombinant soluble IL-6 receptor fused to IL-6, macrophage inflammatory protein 1-alpha (MIP-1-alpha), G-CSF, GM-CSF, thrombopoietin (TPO), platelet factor 4 (PF-4), platelet-derived growth factor (PDGF), neural growth factors and basic fibroblast growth factor (bFGF).

Since totipotent stem cells can give rise to virtually any mature cell type, expansion of these cells in culture will facilitate the production of large quantities of mature cells. Techniques for culturing stem cells are known in the art and administration of polypeptides of the invention, optionally with other growth factors and/or cytokines, is expected to enhance the survival and proliferation of the stem cell populations. This can be accomplished by direct administration of the polypeptide of the invention to the culture medium. Alternatively, stroma cells transfected with a polynucleotide that encodes for the polypeptide of the invention can be used as a feeder layer for the stem cell populations in culture or in vivo. Stromal support cells for feeder layers may include embryonic bone marrow fibroblasts, bone marrow stromal cells, fetal liver cells, or cultured embryonic fibroblasts (see U.S. Patent No. 5,690,926).

Stem cells themselves can be transfected with a polynucleotide of the invention to induce autocrine expression of the polypeptide of the invention. This will allow for generation of undifferentiated totipotential/pluripotential stem cell lines that are useful as is or that can then be differentiated into the desired mature cell types. These stable cell

lines can also serve as a source of undifferentiated totipotential/pluripotential mRNA to create cDNA libraries and templates for polymerase chain reaction experiments. These studies would allow for the isolation and identification of differentially expressed genes in stem cell populations that regulate stem cell proliferation and/or maintenance.

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Expansion and maintenance of totipotent stem cell populations will be useful in the treatment of many pathological conditions. For example, polypeptides of the present invention may be used to manipulate stem cells in culture to give rise to neuroepithelial cells that can be used to augment or replace cells damaged by illness, autoimmune disease, accidental damage or genetic disorders. The polypeptide of the invention may be useful for inducing the proliferation of neural cells and for the regeneration of nerve and brain tissue, i.e. for the treatment of central and peripheral nervous system diseases and neuropathies, as well as mechanical and traumatic disorders which involve degeneration, death or trauma to neural cells or nerve tissue. In addition, the expanded stem cell populations can also be genetically altered for gene therapy purposes and to decrease host rejection of replacement tissues after grafting or implantation.

Expression of the polypeptide of the invention and its effect on stem cells can also be manipulated to achieve controlled differentiation of the stem cells into more differentiated cell types. A broadly applicable method of obtaining pure populations of a specific differentiated cell type from undifferentiated stem cell populations involves the use of a cell-type specific promoter driving a selectable marker. The selectable marker allows only cells of the desired type to survive. For example, stem cells can be induced to differentiate into cardiomyocytes (Wobus et al., Differentiation, 48: 173-182, (1991); Klug et al., J. Clin. Invest., 98(1): 216-224, (1998)) or skeletal muscle cells (Browder, L. W. In: *Principles of Tissue Engineering eds.* Lanza et al., Academic Press (1997)). Alternatively, directed differentiation of stem cells can be accomplished by culturing the

Alternatively, directed differentiation of stem cells can be accomplished by culturing the stem cells in the presence of a differentiation factor such as retinoic acid and an antagonist of the polypeptide of the invention which would inhibit the effects of endogenous stem cell factor activity and allow differentiation to proceed.

In vitro cultures of stem cells can be used to determine if the polypeptide of the invention exhibits stem cell growth factor activity. Stem cells are isolated from any one of various cell sources (including hematopoietic stem cells and embryonic stem cells) and

cultured on a feeder layer, as described by Thompson et al. Proc. Natl. Acad. Sci, U.S.A., 92: 7844-7848 (1995), in the presence of the polypeptide of the invention alone or in combination with other growth factors or cytokines. The ability of the polypeptide of the invention to induce stem cells proliferation is determined by colony formation on semi-solid support e.g. as described by Bernstein et al., Blood, 77: 2316-2321 (1991).

4.10.5 HEMATOPOIESIS REGULATING ACTIVITY

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A polypeptide of the present invention may be involved in regulation of hematopoiesis and, consequently, in the treatment of myeloid or lymphoid cell disorders. 10 Even marginal biological activity in support of colony forming cells or of factor-dependent cell lines indicates involvement in regulating hematopoiesis, e.g. in supporting the growth and proliferation of erythroid progenitor cells alone or in combination with other cytokines, thereby indicating utility, for example, in treating various anemias or for use in conjunction with irradiation/chemotherapy to stimulate the 15 production of erythroid precursors and/or erythroid cells; in supporting the growth and proliferation of myeloid cells such as granulocytes and monocytes/macrophages (i.e., traditional CSF activity) useful, for example, in conjunction with chemotherapy to prevent or treat consequent myelo-suppression; in supporting the growth and proliferation of megakaryocytes and consequently of platelets thereby allowing prevention or treatment of various platelet disorders such as thrombocytopenia, and generally for use in place of or complimentary to platelet transfusions; and/or in supporting the growth and proliferation of hematopoietic stem cells which are capable of maturing to any and all of the above-mentioned hematopoietic cells and therefore find therapeutic utility in various stem cell disorders (such as those usually treated with transplantation, including, without 25 limitation, aplastic anemia and paroxysmal nocturnal hemoglobinuria), as well as in repopulating the stem cell compartment post irradiation/chemotherapy, either in-vivo or ex-vivo (i.e., in conjunction with bone marrow transplantation or with peripheral progenitor cell transplantation (homologous or heterologous)) as normal cells or genetically manipulated for gene therapy.

Therapeutic compositions of the invention can be used in the following:

Suitable assays for proliferation and differentiation of various hematopoietic lines are cited above.

Assays for embryonic stem cell differentiation (which will identify, among others, proteins that influence embryonic differentiation hematopoiesis) include, without limitation, those described in: Johansson et al. Cellular Biology 15:141-151, 1995; Keller et al., Molecular and Cellular Biology 13:473-486, 1993; McClanahan et al., Blood 81:2903-2915, 1993.

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Assays for stem cell survival and differentiation (which will identify, among others, proteins that regulate lympho-hematopoiesis) include, without limitation, those 10 described in: Methylcellulose colony forming assays, Freshney, M. G. In Culture of Hematopoietic Cells. R. I. Freshney, et al. eds. Vol pp. 265-268, Wiley-Liss, Inc., New York, N.Y. 1994; Hirayama et al., Proc. Natl. Acad. Sci. USA 89:5907-5911, 1992; Primitive hematopoietic colony forming cells with high proliferative potential, McNiece, I. K. and Briddell, R. A. In Culture of Hematopoietic Cells. R. I. Freshney, et al. eds. Vol 15 pp. 23-39, Wiley-Liss, Inc., New York, N.Y. 1994; Neben et al., Experimental Hematology 22:353-359, 1994; Cobblestone area forming cell assay, Ploemacher, R. E. In Culture of Hematopoietic Cells. R. I. Freshney, et al. eds. Vol pp. 1-21, Wiley-Liss, Inc., New York, N.Y. 1994; Long term bone marrow cultures in the presence of stromal cells, Spooncer, E., Dexter, M. and Allen, T. In Culture of Hematopoietic Cells. R. L. 20 Freshney, et al. eds. Vol pp. 163-179, Wiley-Liss, Inc., New York, N.Y. 1994; Long term culture initiating cell assay, Sutherland, H. J. In Culture of Hematopoietic Cells. R. I. Freshney, et al. eds. Vol pp. 139-162, Wiley-Liss, Inc., New York, N.Y. 1994.

4.10.6 TISSUE GROWTH ACTIVITY

A polypeptide of the present invention also may be involved in bone, cartilage, tendon, ligament and/or nerve tissue growth or regeneration, as well as in wound healing and tissue repair and replacement, and in healing of burns, incisions and ulcers.

A polypeptide of the present invention which induces cartilage and/or bone growth in circumstances where bone is not normally formed, has application in the healing of bone fractures and cartilage damage or defects in humans and other animals. Compositions of a polypeptide, antibody, binding partner, or other modulator of the

invention may have prophylactic use in closed as well as open fracture reduction and also in the improved fixation of artificial joints. De novo bone formation induced by an osteogenic agent contributes to the repair of congenital, trauma induced, or oncologic resection induced craniofacial defects, and also is useful in cosmetic plastic surgery.

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A polypeptide of this invention may also be involved in attracting bone-forming cells, stimulating growth of bone-forming cells, or inducing differentiation of progenitors of bone-forming cells. Treatment of osteoporosis, osteoarthritis, bone degenerative disorders, or periodontal disease, such as through stimulation of bone and/or cartilage repair or by blocking inflammation or processes of tissue destruction (collagenase activity, osteoclast activity, etc.) mediated by inflammatory processes may also be possible using the composition of the invention.

Another category of tissue regeneration activity that may involve the polypeptide of the present invention is tendon/ligament formation. Induction of tendon/ligament-like tissue or other tissue formation in circumstances where such tissue is not normally formed, has application in the healing of tendon or ligament tears, deformities and other 15 tendon or ligament defects in humans and other animals. Such a preparation employing a tendon/ligament-like tissue inducing protein may have prophylactic use in preventing damage to tendon or ligament tissue, as well as use in the improved fixation of tendon or ligament to bone or other tissues, and in repairing defects to tendon or ligament tissue. De novo tendon/ligament-like tissue formation induced by a composition of the present invention contributes to the repair of congenital, trauma induced, or other tendon or ligament defects of other origin, and is also useful in cosmetic plastic surgery for attachment or repair of tendons or ligaments. The compositions of the present invention may provide environment to attract tendon- or ligament-forming cells, stimulate growth 25 of tendon- or ligament-forming cells, induce differentiation of progenitors of tendon- or ligament-forming cells, or induce growth of tendon/ligament cells or progenitors ex vivo for return in vivo to effect tissue repair. The compositions of the invention may also be useful in the treatment of tendinitis, carpal tunnel syndrome and other tendon or ligament defects. The compositions may also include an appropriate matrix and/or sequestering 30 agent as a carrier as is well known in the art.

The compositions of the present invention may also be useful for proliferation of neural cells and for regeneration of nerve and brain tissue, i.e. for the treatment of central and peripheral nervous system diseases and neuropathies, as well as mechanical and traumatic disorders, which involve degeneration, death or trauma to neural cells or nerve tissue. More specifically, a composition may be used in the treatment of diseases of the peripheral nervous system, such as peripheral nerve injuries, peripheral neuropathy and localized neuropathies, and central nervous system diseases, such as Alzheimer's, Parkinson's disease, Huntington's disease, amyotrophic lateral sclerosis, and Shy-Drager syndrome. Further conditions which may be treated in accordance with the present invention include mechanical and traumatic disorders, such as spinal cord disorders, head trauma and cerebrovascular diseases such as stroke. Peripheral neuropathies resulting from chemotherapy or other medical therapies may also be treatable using a composition of the invention.

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Compositions of the invention may also be useful to promote better or faster closure of non-healing wounds, including without limitation pressure ulcers, ulcers associated with vascular insufficiency, surgical and traumatic wounds, and the like.

Compositions of the present invention may also be involved in the generation or regeneration of other tissues, such as organs (including, for example, pancreas, liver, intestine, kidney, skin, endothelium), muscle (smooth, skeletal or cardiac) and vascular (including vascular endothelium) tissue, or for promoting the growth of cells comprising such tissues. Part of the desired effects may be by inhibition or modulation of fibrotic scarring may allow normal tissue to regenerate. A polypeptide of the present invention may also exhibit angiogenic activity.

A composition of the present invention may also be useful for gut protection or regeneration and treatment of lung or liver fibrosis, reperfusion injury in various tissues, and conditions resulting from systemic cytokine damage.

A composition of the present invention may also be useful for promoting or inhibiting differentiation of tissues described above from precursor tissues or cells; or for inhibiting the growth of tissues described above.

Therapeutic compositions of the invention can be used in the following:

Assays for tissue generation activity include, without limitation, those described in: International Patent Publication No. WO95/16035 (bone, cartilage, tendon); International Patent Publication No. WO95/05846 (nerve, neuronal); International Patent Publication No. WO91/07491 (skin, endothelium).

Assays for wound healing activity include, without limitation, those described in: Winter, Epidermal Wound Healing, pps. 71-112 (Maibach, H. I. and Rovee, D. T., eds.), Year Book Medical Publishers, Inc., Chicago, as modified by Eaglstein and Mertz, J. Invest. Dermatol 71:382-84 (1978).

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4.10.7 IMMUNE STIMULATING OR SUPPRESSING ACTIVITY

A polypeptide of the present invention may also exhibit immune stimulating or immune suppressing activity, including without limitation the activities for which assays are described herein. A polynucleotide of the invention can encode a polypeptide exhibiting such activities. A protein may be useful in the treatment of various immune deficiencies and disorders (including severe combined immunodeficiency (SCID)), e.g., in regulating (up or down) growth and proliferation of T and/or B lymphocytes, as well as effecting the cytolytic activity of NK cells and other cell populations. These immune deficiencies may be genetic or be caused by viral (e.g., HIV) as well as bacterial or fungal infections, or may result from autoimmune disorders. More specifically, infectious diseases causes by viral, bacterial, fungal or other infection may be treatable using a protein of the present invention, including infections by HIV, hepatitis viruses, herpes viruses, mycobacteria, Leishmania spp., malaria spp. and various fungal infections such as candidiasis. Of course, in this regard, proteins of the present invention may also be useful where a boost to the immune system generally may be desirable, i.e., in the treatment of cancer.

Autoimmune disorders which may be treated using a protein of the present invention include, for example, connective tissue disease, multiple sclerosis, systemic lupus erythematosus, rheumatoid arthritis, autoimmune pulmonary inflammation, Guillain-Barre syndrome, autoimmune thyroiditis, insulin dependent diabetes mellitis, myasthenia gravis, graft-versus-host disease and autoimmune inflammatory eye disease. Such a protein (or antagonists thereof, including antibodies) of the present invention may

also to be useful in the treatment of allergic reactions and conditions (e.g., anaphylaxis, serum sickness, drug reactions, food allergies, insect venom allergies, mastocytosis, allergic rhinitis, hypersensitivity pneumonitis, urticaria, angioedema, eczema, atopic dermatitis, allergic contact dermatitis, erythema multiforme, Stevens-Johnson syndrome, allergic conjunctivitis, atopic keratoconjunctivitis, venereal keratoconjunctivitis, giant papillary conjunctivitis and contact allergies), such as asthma (particularly allergic asthma) or other respiratory problems. Other conditions, in which immune suppression is desired (including, for example, organ transplantation), may also be treatable using a protein (or antagonists thereof) of the present invention. The therapeutic effects of the polypeptides or antagonists thereof on allergic reactions can be evaluated by in vivo animals models such as the cumulative contact enhancement test (Lastbom et al., Toxicology 125: 59-66, 1998), skin prick test (Hoffmann et al., Allergy 54: 446-54, 1999), guinea pig skin sensitization test (Vohr et al., Arch. Toxocol. 73: 501-9), and murine local lymph node assay (Kimber et al., J. Toxicol. Environ. Health 53: 563-79).

Using the proteins of the invention it may also be possible to modulate immune responses, in a number of ways. Down regulation may be in the form of inhibiting or blocking an immune response already in progress or may involve preventing the induction of an immune response. The functions of activated T cells may be inhibited by suppressing T cell responses or by inducing specific tolerance in T cells, or both. Immunosuppression of T cell responses is generally an active, non-antigen-specific, process which requires continuous exposure of the T cells to the suppressive agent. Tolerance, which involves inducing non-responsiveness or anergy in T cells, is distinguishable from immunosuppression in that it is generally antigen-specific and persists after exposure to the tolerizing agent has ceased. Operationally, tolerance can be demonstrated by the lack of a T cell response upon reexposure to specific antigen in the absence of the tolerizing agent.

Down regulating or preventing one or more antigen functions (including without limitation B lymphocyte antigen functions (such as, for example, B7)), e.g., preventing high level lymphokine synthesis by activated T cells, will be useful in situations of tissue, skin and organ transplantation and in graft-versus-host disease (GVHD). For example, blockage of T cell function should result in reduced tissue destruction in tissue

transplantation. Typically, in tissue transplants, rejection of the transplant is initiated through its recognition as foreign by T cells, followed by an immune reaction that destroys the transplant. The administration of a therapeutic composition of the invention may prevent cytokine synthesis by immune cells, such as T cells, and thus acts as an immunosuppressant. Moreover, a lack of costimulation may also be sufficient to anergize the T cells, thereby inducing tolerance in a subject. Induction of long-term tolerance by B lymphocyte antigen-blocking reagents may avoid the necessity of repeated administration of these blocking reagents. To achieve sufficient immunosuppression or tolerance in a subject, it may also be necessary to block the function of a combination of B lymphocyte antigens.

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The efficacy of particular therapeutic compositions in preventing organ transplant rejection or GVHD can be assessed using animal models that are predictive of efficacy in humans. Examples of appropriate systems which can be used include allogeneic cardiac grafts in rats and xenogeneic pancreatic islet cell grafts in mice, both of which have been used to examine the immunosuppressive effects of CTLA4Ig fusion proteins in vivo as described in Lenschow et al., Science 257:789-792 (1992) and Turka et al., Proc. Natl. Acad. Sci USA, 89:11102-11105 (1992). In addition, murine models of GVHD (see Paul ed., Fundamental Immunology, Raven Press, New York, 1989, pp. 846-847) can be used to determine the effect of therapeutic compositions of the invention on the development of that disease.

Blocking antigen function may also be therapeutically useful for treating autoimmune diseases. Many autoimmune disorders are the result of inappropriate activation of T cells that are reactive against self tissue and which promote the production of cytokines and autoantibodies involved in the pathology of the diseases. Preventing the activation of autoreactive T cells may reduce or eliminate disease symptoms.

Administration of reagents which block stimulation of T cells can be used to inhibit T cell activation and prevent production of autoantibodies or T cell-derived cytokines which may be involved in the disease process. Additionally, blocking reagents may induce antigen-specific tolerance of autoreactive T cells which could lead to long-term relief from the disease. The efficacy of blocking reagents in preventing or alleviating autoimmune disorders can be determined using a number of well-characterized animal

models of human autoimmune diseases. Examples include murine experimental autoimmune encephalitis, systemic lupus erythmatosis in MRL/lpr/lpr mice or NZB hybrid mice, murine autoimmune collagen arthritis, diabetes mellitus in NOD mice and BB rats, and murine experimental myasthenia gravis (see Paul ed., Fundamental Immunology, Raven Press, New York, 1989, pp. 840-856).

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Upregulation of an antigen function (e.g., a B lymphocyte antigen function), as a means of up regulating immune responses, may also be useful in therapy. Upregulation of immune responses may be in the form of enhancing an existing immune response or eliciting an initial immune response. For example, enhancing an immune response may be useful in cases of viral infection, including systemic viral diseases such as influenza, the common cold, and encephalitis.

Alternatively, anti-viral immune responses may be enhanced in an infected patient by removing T cells from the patient, costimulating the T cells in vitro with viral antigen-pulsed APCs either expressing a peptide of the present invention or together with a stimulatory form of a soluble peptide of the present invention and reintroducing the in vitro activated T cells into the patient. Another method of enhancing anti-viral immune responses would be to isolate infected cells from a patient, transfect them with a nucleic acid encoding a protein of the present invention as described herein such that the cells express all or a portion of the protein on their surface, and reintroduce the transfected cells into the patient. The infected cells would now be capable of delivering a costimulatory signal to, and thereby activate, T cells in vivo.

A polypeptide of the present invention may provide the necessary stimulation signal to T cells to induce a T cell mediated immune response against the transfected tumor cells. In addition, tumor cells which lack MHC class I or MHC class II molecules, or which fail to reexpress sufficient mounts of MHC class I or MHC class II molecules, can be transfected with nucleic acid encoding all or a portion of (e.g., a cytoplasmic-domain truncated portion) of an MHC class I alpha chain protein and β_2 microglobulin protein or an MHC class II alpha chain protein and an MHC class II beta chain protein to thereby express MHC class I or MHC class II proteins on the cell surface. Expression of the appropriate class I or class II MHC in conjunction with a peptide having the activity of a B lymphocyte antigen (e.g., B7-1, B7-2, B7-3) induces a

T cell mediated immune response against the transfected tumor cell. Optionally, a gene encoding an antisense construct which blocks expression of an MHC class II associated protein, such as the invariant chain, can also be cotransfected with a DNA encoding a peptide having the activity of a B lymphocyte antigen to promote presentation of tumor associated antigens and induce tumor specific immunity. Thus, the induction of a T cell mediated immune response in a human subject may be sufficient to overcome tumor-specific tolerance in the subject.

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The activity of a protein of the invention may, among other means, be measured by the following methods:

Suitable assays for thymocyte or splenocyte cytotoxicity include, without limitation, those described in: Current Protocols in Immunology, Ed by J. E. Coligan, A. M. Kruisbeek, D. H. Margulies, E. M. Shevach, W. Strober, Pub. Greene Publishing Associates and Wiley-Interscience (Chapter 3, In Vitro assays for Mouse Lymphocyte Function 3.1-3.19; Chapter 7, Immunologic studies in Humans); Herrmann et al., Proc. Natl. Acad. Sci. USA 78:2488-2492, 1981; Herrmann et al., J. Immunol. 128:1968-1974, 1982; Handa et al., J. Immunol. 135:1564-1572, 1985; Takai et al., I. Immunol. 137:3494-3500, 1986; Takai et al., J. Immunol. 140:508-512, 1988; Bowman et al., J. Virology 61:1992-1998; Bertagnolli et al., Cellular Immunology 133:327-341, 1991; Brown et al., J. Immunol. 153:3079-3092, 1994.

Assays for T-cell-dependent immunoglobulin responses and isotype switching (which will identify, among others, proteins that modulate T-cell dependent antibody responses and that affect Th1/Th2 profiles) include, without limitation, those described in: Maliszewski, J. Immunol. 144:3028-3033, 1990; and Assays for B cell function: In vitro antibody production, Mond, J. J. and Brunswick, M. In Current Protocols in Immunology. J. E. e.a. Coligan eds. Vol 1 pp. 3.8.1-3.8.16, John Wiley and Sons, Toronto, 1994.

Mixed lymphocyte reaction (MLR) assays (which will identify, among others, proteins that generate predominantly Th1 and CTL responses) include, without limitation, those described in: Current Protocols in Immunology, Ed by J. E. Coligan, A. M. Kruisbeek, D. H. Margulies, E. M. Shevach, W. Strober, Pub. Greene Publishing Associates and Wiley-Interscience (Chapter 3, In Vitro assays for Mouse Lymphocyte

Function 3.1-3.19; Chapter 7, Immunologic studies in Humans); Takai et al., J. Immunol. 137:3494-3500, 1986; Takai et al., J. Immunol. 140:508-512, 1988; Bertagnolli et al., J. Immunol. 149:3778-3783, 1992.

Dendritic cell-dependent assays (which will identify, among others, proteins expressed by dendritic cells that activate naive T-cells) include, without limitation, those described in: Guery et al., J. Immunol. 134:536-544, 1995; Inaba et al., Journal of Experimental Medicine 173:549-559, 1991; Macatonia et al., Journal of Immunology 154:5071-5079, 1995; Porgador et al., Journal of Experimental Medicine 182:255-260, 1995; Nair et al., Journal of Virology 67:4062-4069, 1993; Huang et al., Science 264:961-965, 1994; Macatonia et al., Journal of Experimental Medicine 169:1255-1264, 1989; Bhardwaj et al., Journal of Clinical Investigation 94:797-807, 1994; and Inaba et al., Journal of Experimental Medicine 172:631-640, 1990.

Assays for lymphocyte survival/apoptosis (which will identify, among others, proteins that prevent apoptosis after superantigen induction and proteins that regulate lymphocyte homeostasis) include, without limitation, those described in: Darzynkiewicz et al., Cytometry 13:795-808, 1992; Gorczyca et al., Leukemia 7:659-670, 1993; Gorczyca et al., Cancer Research 53:1945-1951, 1993; Itoh et al., Cell 66:233-243, 1991; Zacharchuk, Journal of Immunology 145:4037-4045, 1990; Zamai et al., Cytometry 14:891-897, 1993; Gorczyca et al., International Journal of Oncology 1:639-648, 1992.

Assays for proteins that influence early steps of T-cell commitment and development include, without limitation, those described in: Antica et al., Blood 84:111-117, 1994; Fine et al., Cellular Immunology 155:111-122, 1994; Galy et al., Blood 85:2770-2778, 1995; Toki et al., Proc. Nat. Acad Sci. USA 88:7548-7551, 1991.

25 4.10.8 ACTIVIN/INHIBIN ACTIVITY

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A polypeptide of the present invention may also exhibit activin- or inhibin-related activities. A polynucleotide of the invention may encode a polypeptide exhibiting such characteristics. Inhibins are characterized by their ability to inhibit the release of follicle stimulating hormone (FSH), while activins and are characterized by their ability to stimulate the release of follicle stimulating hormone (FSH). Thus, a polypeptide of the present invention, alone or in heterodimers with a member of the inhibin family, may be

useful as a contraceptive based on the ability of inhibins to decrease fertility in female mammals and decrease spermatogenesis in male mammals. Administration of sufficient amounts of other inhibins can induce infertility in these mammals. Alternatively, the polypeptide of the invention, as a homodimer or as a heterodimer with other protein subunits of the inhibin group, may be useful as a fertility inducing therapeutic, based upon the ability of activin molecules in stimulating FSH release from cells of the anterior pituitary. See, for example, U.S. Pat. No. 4,798,885. A polypeptide of the invention may also be useful for advancement of the onset of fertility in sexually immature mammals, so as to increase the lifetime reproductive performance of domestic animals such as, but not limited to, cows, sheep and pigs.

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The activity of a polypeptide of the invention may, among other means, be measured by the following methods.

Assays for activin/inhibin activity include, without limitation, those described in: Vale et al., Endocrinology 91:562-572, 1972; Ling et al., Nature 321:779-782, 1986; Vale et al., Nature 321:776-779, 1986; Mason et al., Nature 318:659-663, 1985; Forage et al., Proc. Natl. Acad. Sci. USA 83:3091-3095, 1986.

4.10.9 CHEMOTACTIC/CHEMOKINETIC ACTIVITY

A polypeptide of the present invention may be involved in chemotactic or chemokinetic activity for mammalian cells, including, for example, monocytes, fibroblasts, neutrophils, T-cells, mast cells, eosinophils, epithelial and/or endothelial cells. A polynucleotide of the invention can encode a polypeptide exhibiting such attributes. Chemotactic and chemokinetic receptor activation can be used to mobilize or attract a desired cell population to a desired site of action. Chemotactic or chemokinetic compositions (e.g. proteins, antibodies, binding partners, or modulators of the invention) provide particular advantages in treatment of wounds and other trauma to tissues, as well as in treatment of localized infections. For example, attraction of lymphocytes, monocytes or neutrophils to tumors or sites of infection may result in improved immune responses against the tumor or infecting agent.

A protein or peptide has chemotactic activity for a particular cell population if it can stimulate, directly or indirectly, the directed orientation or movement of such cell

population. Preferably, the protein or peptide has the ability to directly stimulate directed movement of cells. Whether a particular protein has chemotactic activity for a population of cells can be readily determined by employing such protein or peptide in any known assay for cell chemotaxis.

5 Therapeutic compositions of the invention can be used in the following:

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Assays for chemotactic activity (which will identify proteins that induce or prevent chemotaxis) consist of assays that measure the ability of a protein to induce the migration of cells across a membrane as well as the ability of a protein to induce the adhesion of one cell population to another cell population. Suitable assays for movement and adhesion include, without limitation, those described in: Current Protocols in Immunology, Ed by J. E. Coligan, A. M. Kruisbeek, D. H. Marguiles, E. M. Shevach, W. Strober, Pub. Greene Publishing Associates and Wiley-Interscience (Chapter 6.12, Measurement of alpha and beta Chemokines 6.12.1-6.12.28; Taub et al. J. Clin. Invest. 95:1370-1376, 1995; Lind et al. APMIS 103:140-146, 1995; Muller et al Eur. J. Immunol. 25:1744-1748; Gruber et al. J. of Immunol. 152:5860-5867, 1994; Johnston et al. J. of Immunol. 153:1762-1768, 1994.

4.10.10 HEMOSTATIC AND THROMBOLYTIC ACTIVITY

A polypeptide of the invention may also be involved in hemostatis or

thrombolysis or thrombosis. A polynucleotide of the invention can encode a polypeptide exhibiting such attributes. Compositions may be useful in treatment of various coagulation disorders (including hereditary disorders, such as hemophilias) or to enhance coagulation and other hemostatic events in treating wounds resulting from trauma, surgery or other causes. A composition of the invention may also be useful for dissolving or inhibiting formation of thromboses and for treatment and prevention of conditions resulting therefrom (such as, for example, infarction of cardiac and central nervous system vessels (e.g., stroke).

Therapeutic compositions of the invention can be used in the following:

Assay for hemostatic and thrombolytic activity include, without limitation, those described in: Linet et al., J. Clin. Pharmacol. 26:131-140, 1986; Burdick et al.,

Thrombosis Res. 45:413-419, 1987; Humphrey et al., Fibrinolysis 5:71-79 (1991); Schaub, Prostaglandins 35:467-474, 1988.

4.10.11 CANCER DIAGNOSIS AND THERAPY

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Polypeptides of the invention may be involved in cancer cell generation, proliferation or metastasis. Detection of the presence or amount of polynucleotides or polypeptides of the invention may be useful for the diagnosis and/or prognosis of one or more types of cancer. For example, the presence or increased expression of a polynucleotide/polypeptide of the invention may indicate a hereditary risk of cancer, a precancerous condition, or an ongoing malignancy. Conversely, a defect in the gene or absence of the polypeptide may be associated with a cancer condition. Identification of single nucleotide polymorphisms associated with cancer or a predisposition to cancer may also be useful for diagnosis or prognosis.

Cancer treatments promote tumor regression by inhibiting tumor cell proliferation, inhibiting angiogenesis (growth of new blood vessels that is necessary to support tumor growth) and/or prohibiting metastasis by reducing tumor cell motility or invasiveness. Therapeutic compositions of the invention may be effective in adult and pediatric oncology including in solid phase tumors/malignancies, locally advanced tumors, human soft tissue sarcomas, metastatic cancer, including lymphatic metastases, blood cell malignancies including multiple myeloma, acute and chronic leukemias, and lymphomas, head and neck cancers including mouth cancer, larynx cancer and thyroid cancer, lung cancers including small cell carcinoma and non-small cell cancers, breast cancers including small cell carcinoma and ductal carcinoma, gastrointestinal cancers including esophageal cancer, stomach cancer, colon cancer, colorectal cancer and polyps associated with colorectal neoplasia, pancreatic cancers, liver cancer, urologic cancers including bladder cancer and prostate cancer, malignancies of the female genital tract including ovarian carcinoma, uterine (including endometrial) cancers, and solid tumor in the ovarian follicle, kidney cancers including renal cell carcinoma, brain cancers including intrinsic brain tumors, neuroblastoma, astrocytic brain tumors, gliomas, metastatic tumor cell invasion in the central nervous system, bone cancers including osteomas, skin cancers including malignant melanoma, tumor progression of human skin

keratinocytes, squamous cell carcinoma, basal cell carcinoma, hemangiopericytoma and Karposi's sarcoma.

Polypeptides, polynucleotides, or modulators of polypeptides of the invention (including inhibitors and stimulators of the biological activity of the polypeptide of the invention) may be administered to treat cancer. Therapeutic compositions can be administered in therapeutically effective dosages alone or in combination with adjuvant cancer therapy such as surgery, chemotherapy, radiotherapy, thermotherapy, and laser therapy, and may provide a beneficial effect, e.g. reducing tumor size, slowing rate of tumor growth, inhibiting metastasis, or otherwise improving overall clinical condition, without necessarily eradicating the cancer.

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The composition can also be administered in therapeutically effective amounts as a portion of an anti-cancer cocktail. An anti-cancer cocktail is a mixture of the polypeptide or modulator of the invention with one or more anti-cancer drugs in addition to a pharmaceutically acceptable carrier for delivery. The use of anti-cancer cocktails as 15 a cancer treatment is routine. Anti-cancer drugs that are well known in the art and can be used as a treatment in combination with the polypeptide or modulator of the invention include: Actinomycin D, Aminoglutethimide, Asparaginase, Bleomycin, Busulfan, Carboplatin, Carmustine, Chlorambucil, Cisplatin (cis-DDP), Cyclophosphamide, Cytarabine HCl (Cytosine arabinoside), Dacarbazine, Dactinomycin, Daunorubicin HCl, Doxorubicin HCl, Estramustine phosphate sodium, Etoposide (V16-213), Floxuridine, 5-20. Fluorouracil (5-Fu), Flutamide, Hydroxyurea (hydroxycarbamide), Ifosfamide, Interferon Alpha-2a, Interferon Alpha-2b, Leuprolide acetate (LHRH-releasing factor analog), Lomustine, Mechlorethamine HCl (nitrogen mustard), Melphalan, Mercaptopurine, Mesna, Methotrexate (MTX), Mitomycin, Mitoxantrone HCl, Octreotide, Plicamycin, 25 Procarbazine HCl, Streptozocin, Tamoxifen citrate, Thioguanine, Thiotepa, Vinblastine sulfate, Vincristine sulfate, Amsacrine, Azacitidine, Hexamethylmelamine, Interleukin-2, Mitoguazone, Pentostatin, Semustine, Teniposide, and Vindesine sulfate.

In addition, therapeutic compositions of the invention may be used for prophylactic treatment of cancer. There are hereditary conditions and/or environmental situations (e.g. exposure to carcinogens) known in the art that predispose an individual to developing cancers. Under these circumstances, it may be beneficial to treat these

individuals with therapeutically effective doses of the polypeptide of the invention to reduce the risk of developing cancers.

In vitro models can be used to determine the effective doses of the polypeptide of the invention as a potential cancer treatment. These in vitro models include proliferation assays of cultured tumor cells, growth of cultured tumor cells in soft agar (see Freshney, (1987) Culture of Animal Cells: A Manual of Basic Technique, Wily-Liss, New York, NY Ch 18 and Ch 21), tumor systems in nude mice as described in Giovanella et al., J. Natl. Can. Inst., 52: 921-30 (1974), mobility and invasive potential of tumor cells in Boyden Chamber assays as described in Pilkington et al., Anticancer Res., 17: 4107-9 (1997), and angiogenesis assays such as induction of vascularization of the chick chorioallantoic membrane or induction of vascular endothelial cell migration as described in Ribatta et al., Intl. J. Dev. Biol., 40: 1189-97 (1999) and Li et al., Clin. Exp. Metastasis, 17:423-9 (1999), respectively. Suitable tumor cells lines are available, e.g. from American Type Tissue Culture Collection catalogs.

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4.10.12 RECEPTOR/LIGAND ACTIVITY

A polypeptide of the present invention may also demonstrate activity as receptor, receptor ligand or inhibitor or agonist of receptor/ligand interactions. A polynucleotide of the invention can encode a polypeptide exhibiting such characteristics. Examples of such receptors and ligands include, without limitation, cytokine receptors and their ligands, receptor kinases and their ligands, receptor phosphatases and their ligands, receptors involved in cell-cell interactions and their ligands (including without limitation, cellular adhesion molecules (such as selectins, integrins and their ligands) and receptor/ligand pairs involved in antigen presentation, antigen recognition and development of cellular and humoral immune responses. Receptors and ligands are also useful for screening of potential peptide or small molecule inhibitors of the relevant receptor/ligand interaction. A protein of the present invention (including, without limitation, fragments of receptors and ligands) may themselves be useful as inhibitors of receptor/ligand interactions.

The activity of a polypeptide of the invention may, among other means, be measured by the following methods:

Suitable assays for receptor-ligand activity include without limitation those described in: Current Protocols in Immunology, Ed by J. E. Coligan, A. M. Kruisbeek, D. H. Margulies, E. M. Shevach, W. Strober, Pub. Greene Publishing Associates and Wiley- Interscience (Chapter 7.28, Measurement of Cellular Adhesion under static conditions 7.28.1-7.28.22), Takai et al., Proc. Natl. Acad. Sci. USA 84:6864-6868, 1987; Bierer et al., J. Exp. Med. 168:1145-1156, 1988; Rosenstein et al., J. Exp. Med. 169:149-160 1989; Stoltenborg et al., J. Immunol. Methods 175:59-68, 1994; Stitt et al., Cell 80:661-670, 1995.

By way of example, the polypeptides of the invention may be used as a receptor for a ligand(s) thereby transmitting the biological activity of that ligand(s). Ligands may be identified through binding assays, affinity chromatography, dihybrid screening assays, BIAcore assays, gel overlay assays, or other methods known in the art.

Studies characterizing drugs or proteins as agonist or antagonist or partial agonists or a partial antagonist require the use of other proteins as competing ligands. The polypeptides of the present invention or ligand(s) thereof may be labeled by being coupled to radioisotopes, colorimetric molecules or a toxin molecules by conventional methods. ("Guide to Protein Purification" Murray P. Deutscher (ed) Methods in Enzymology Vol. 182 (1990) Academic Press, Inc. San Diego). Examples of radioisotopes include, but are not limited to, tritium and carbon-14. Examples of colorimetric molecules include, but are not limited to, fluorescent molecules such as fluorescamine, or rhodamine or other colorimetric molecules. Examples of toxins include, but are not limited, to ricin.

4.10.13 DRUG SCREENING

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This invention is particularly useful for screening chemical compounds by using the novel polypeptides or binding fragments thereof in any of a variety of drug screening techniques. The polypeptides or fragments employed in such a test may either be free in solution, affixed to a solid support, borne on a cell surface or located intracellularly. One method of drug screening utilizes eukaryotic or prokaryotic host cells which are stably transformed with recombinant nucleic acids expressing the polypeptide or a fragment thereof. Drugs are screened against such transformed cells in competitive binding assays.

Such cells, either in viable or fixed form, can be used for standard binding assays. One may measure, for example, the formation of complexes between polypeptides of the invention or fragments and the agent being tested or examine the diminution in complex formation between the novel polypeptides and an appropriate cell line, which are well known in the art.

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Sources for test compounds that may be screened for ability to bind to or modulate (i.e., increase or decrease) the activity of polypeptides of the invention include (1) inorganic and organic chemical libraries, (2) natural product libraries, and (3) combinatorial libraries comprised of either random or mimetic peptides, oligonucleotides or organic molecules.

Chemical libraries may be readily synthesized or purchased from a number of commercial sources, and may include structural analogs of known compounds or compounds that are identified as "hits" or "leads" via natural product screening.

The sources of natural product libraries are microorganisms (including bacteria and fungi), animals, plants or other vegetation, or marine organisms, and libraries of mixtures for screening may be created by: (1) fermentation and extraction of broths from soil, plant or marine microorganisms or (2) extraction of the organisms themselves.

Natural product libraries include polyketides, non-ribosomal peptides, and (non-naturally occurring) variants thereof. For a review, see *Science 282:63-68* (1998).

Combinatorial libraries are composed of large numbers of peptides, oligonucleotides or organic compounds and can be readily prepared by traditional automated synthesis methods, PCR, cloning or proprietary synthetic methods. Of particular interest are peptide and oligonucleotide combinatorial libraries. Still other libraries of interest include peptide, protein, peptidomimetic, multiparallel synthetic collection, recombinatorial, and polypeptide libraries. For a review of combinatorial chemistry and libraries created therefrom, see Myers, Curr. Opin. Biotechnol. 8:701-707 (1997). For reviews and examples of peptidomimetic libraries, see Al-Obeidi et al., Mol. Biotechnol, 9(3):205-23 (1998); Hruby et al., Curr Opin Chem Biol, 1(1):114-19 (1997); Dorner et al., Bioorg Med Chem, 4(5):709-15 (1996) (alkylated dipeptides).

Identification of modulators through use of the various libraries described herein permits modification of the candidate "hit" (or "lead") to optimize the capacity of the

"hit" to bind a polypeptide of the invention. The molecules identified in the binding assay are then tested for antagonist or agonist activity in *in vivo* tissue culture or animal models that are well known in the art. In brief, the molecules are titrated into a plurality of cell cultures or animals and then tested for either cell/animal death or prolonged survival of the animal/cells.

The binding molecules thus identified may be complexed with toxins, e.g., ricin or cholera, or with other compounds that are toxic to cells such as radioisotopes. The toxin-binding molecule complex is then targeted to a tumor or other cell by the specificity of the binding molecule for a polypeptide of the invention. Alternatively, the binding molecules may be complexed with imaging agents for targeting and imaging purposes.

4.10.14 ASSAY FOR RECEPTOR ACTIVITY

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The invention also provides methods to detect specific binding of a polypeptide e.g. a ligand or a receptor. The art provides numerous assays particularly useful for identifying previously unknown binding partners for receptor polypeptides of the invention. For example, expression cloning using mammalian or bacterial cells, or dihybrid screening assays can be used to identify polynucleotides encoding binding partners. As another example, affinity chromatography with the appropriate immobilized polypeptide of the invention can be used to isolate polypeptides that recognize and bind polypeptides of the invention. There are a number of different libraries used for the identification of compounds, and in particular small molecules, that modulate (i.e., increase or decrease) biological activity of a polypeptide of the invention. Ligands for receptor polypeptides of the invention can also be identified by adding exogenous ligands, or cocktails of ligands to two cells populations that are genetically identical except for the expression of the receptor of the invention: one cell population expresses the receptor of the invention whereas the other does not. The response of the two cell populations to the addition of ligands(s) are then compared. Alternatively, an expression library can be co-expressed with the polypeptide of the invention in cells and assayed for an autocrine response to identify potential ligand(s). As still another example, BIAcore assays, gel overlay assays, or other methods known in the art can be used to identify binding partner polypeptides, including, (1) organic and inorganic chemical libraries, (2)

natural product libraries, and (3) combinatorial libraries comprised of random peptides, oligonucleotides or organic molecules.

The role of downstream intracellular signaling molecules in the signaling cascade of the polypeptide of the invention can be determined. For example, a chimeric protein in which the cytoplasmic domain of the polypeptide of the invention is fused to the extracellular portion of a protein, whose ligand has been identified, is produced in a host cell. The cell is then incubated with the ligand specific for the extracellular portion of the chimeric protein, thereby activating the chimeric receptor. Known downstream proteins involved in intracellular signaling can then be assayed for expected modifications i.e. phosphorylation. Other methods known to those in the art can also be used to identify signaling molecules involved in receptor activity.

4.10.15 ANTI-INFLAMMATORY ACTIVITY

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Compositions of the present invention may also exhibit anti-inflammatory 15 activity. The anti-inflammatory activity may be achieved by providing a stimulus to cells involved in the inflammatory response, by inhibiting or promoting cell-cell interactions (such as, for example, cell adhesion), by inhibiting or promoting chemotaxis of cells involved in the inflammatory process, inhibiting or promoting cell extravasation, or by stimulating or suppressing production of other factors which more directly inhibit or 20 promote an inflammatory response. Compositions with such activities can be used to treat inflammatory conditions including chronic or acute conditions), including without limitation intimation associated with infection (such as septic shock, sepsis or systemic inflammatory response syndrome (SIRS)), ischemia-reperfusion injury, endotoxin lethality, arthritis, complement-mediated hyperacute rejection, nephritis, cytokine or 25 chemokine-induced lung injury, inflammatory bowel disease, Crohn's disease or resulting from over production of cytokines such as TNF or IL-1. Compositions of the invention may also be useful to treat anaphylaxis and hypersensitivity to an antigenic substance or material. Compositions of this invention may be utilized to prevent or treat conditions such as, but not limited to, sepsis, acute pancreatitis, endotoxin shock, cytokine induced 30 shock, rheumatoid arthritis, chronic inflammatory arthritis, pancreatic cell damage from diabetes mellitus type 1, graft versus host disease, inflammatory bowel disease,

inflamation associated with pulmonary disease, other autoimmune disease or inflammatory disease, an antiproliferative agent such as for acute or chronic mylegenous leukemia or in the prevention of premature labor secondary to intrauterine infections.

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Leukemias and related disorders may be treated or prevented by administration of a therapeutic that promotes or inhibits function of the polynucleotides and/or polypeptides of the invention. Such leukemias and related disorders include but are not limited to acute leukemia, acute lymphocytic leukemia, acute myelocytic leukemia, myeloblastic, promyelocytic, myelomonocytic, monocytic, erythroleukemia, chronic leukemia, chronic myelocytic (granulocytic) leukemia and chronic lymphocytic leukemia (for a review of such disorders, see Fishman et al., 1985, Medicine, 2d Ed., J.B. Lippincott Co., Philadelphia).

15 4.10.17 NERVOUS SYSTEM DISORDERS

Nervous system disorders, involving cell types which can be tested for efficacy of intervention with compounds that modulate the activity of the polynucleotides and/or polypeptides of the invention, and which can be treated upon thus observing an indication of therapeutic utility, include but are not limited to nervous system injuries, and diseases or disorders which result in either a disconnection of axons, a diminution or degeneration of neurons, or demyelination. Nervous system lesions which may be treated in a patient (including human and non-human mammalian patients) according to the invention include but are not limited to the following lesions of either the central (including spinal cord, brain) or peripheral nervous systems:

- (i) traumatic lesions, including lesions caused by physical injury or associated with surgery, for example, lesions which sever a portion of the nervous system, or compression injuries;
 - (ii) ischemic lesions, in which a lack of oxygen in a portion of the nervous system results in neuronal injury or death, including cerebral infarction or ischemia, or spinal cord infarction or ischemia;

(iii) infectious lesions, in which a portion of the nervous system is destroyed or injured as a result of infection, for example, by an abscess or associated with infection by human immunodeficiency virus, herpes zoster, or herpes simplex virus or with Lyme disease, tuberculosis, syphilis;

(iv) ___degenerative lesions, in which a portion of the nervous system is destroyed or injured as a result of a degenerative process including but not limited to degeneration associated with Parkinson's disease, Alzheimer's disease, Huntington's chorea, or amyotrophic lateral sclerosis;

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- (v) lesions associated with nutritional diseases or disorders, in which a portion of the nervous system is destroyed or injured by a nutritional disorder or disorder of metabolism including but not limited to, vitamin B12 deficiency, folic acid deficiency, Wernicke disease, tobacco-alcohol amblyopia, Marchiafava-Bignami disease (primary degeneration of the corpus callosum), and alcoholic cerebellar degeneration;
 - (vi) neurological lesions associated with systemic diseases including but not limited to diabetes (diabetic neuropathy, Bell's palsy), systemic lupus erythematosus, carcinoma, or sarcoidosis;
 - (vii) lesions caused by toxic substances including alcohol, lead, or particular neurotoxins; and
- (viii) demyelinated lesions in which a portion of the nervous system is destroyed or injured by a demyelinating disease including but not limited to multiple sclerosis, human immunodeficiency virus-associated myelopathy, transverse myelopathy or various etiologies, progressive multifocal leukoencephalopathy, and central pontine myelinolysis.

Therapeutics which are useful according to the invention for treatment of a nervous system disorder may be selected by testing for biological activity in promoting the survival or differentiation of neurons. For example, and not by way of limitation, therapeutics which elicit any of the following effects may be useful according to the invention:

- (i) increased survival time of neurons in culture;
- 30 (ii) increased sprouting of neurons in culture or in vivo;

(iii) increased production of a neuron-associated molecule in culture or *in vivo*, e.g., choline acetyltransferase or acetylcholinesterase with respect to motor neurons; or

(iv) decreased symptoms of neuron dysfunction in vivo.

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Such effects may be measured by any method known in the art. In preferred, non-limiting embodiments, increased survival of neurons may be measured by the method set forth in Arakawa et al. (1990, J. Neurosci. 10:3507-3515); increased sprouting of neurons may be detected by methods set forth in Pestronk et al. (1980, Exp. Neurol. 70:65-82) or Brown et al. (1981, Ann. Rev. Neurosci. 4:17-42); increased production of neuron-associated molecules may be measured by bioassay, enzymatic assay, antibody binding, Northern blot assay, etc., depending on the molecule to be measured; and motor neuron dysfunction may be measured by assessing the physical manifestation of motor neuron disorder, e.g., weakness, motor neuron conduction velocity, or functional disability.

In specific embodiments, motor neuron disorders that may be treated according to the invention include but are not limited to disorders such as infarction, infection, exposure to toxin, trauma, surgical damage, degenerative disease or malignancy that may affect motor neurons as well as other components of the nervous system, as well as disorders that selectively affect neurons such as amyotrophic lateral sclerosis, and including but not limited to progressive spinal muscular atrophy, progressive bulbar palsy, primary lateral sclerosis, infantile and juvenile muscular atrophy, progressive bulbar paralysis of childhood (Fazio-Londe syndrome), poliomyelitis and the post polio syndrome, and Hereditary Motorsensory Neuropathy (Charcot-Marie-Tooth Disease).

4.10.18 OTHER ACTIVITIES

A polypeptide of the invention may also exhibit one or more of the following additional activities or effects: inhibiting the growth, infection or function of, or killing, infectious agents, including, without limitation, bacteria, viruses, fungi and other parasites; effecting (suppressing or enhancing) bodily characteristics, including, without limitation, height, weight, hair color, eye color, skin, fat to lean ratio or other tissue pigmentation, or organ or body part size or shape (such as, for example, breast augmentation or diminution, change in bone form or shape); effecting biorhythms or

circadian cycles or rhythms; effecting the fertility of male or female subjects; effecting the metabolism, catabolism, anabolism, processing, utilization, storage or elimination of dietary fat, lipid, protein, carbohydrate, vitamins, minerals, co-factors or other nutritional factors or component(s); effecting behavioral characteristics, including, without limitation, appetite, libido, stress, cognition (including cognitive disorders), depression (including depressive disorders) and violent behaviors; providing analgesic effects or other pain reducing effects; promoting differentiation and growth of embryonic stem cells in lineages other than hematopoietic lineages; hormonal or endocrine activity; in the case of enzymes, correcting deficiencies of the enzyme and treating deficiency-related diseases; treatment of hyperproliferative disorders (such as, for example, psoriasis); immunoglobulin-like activity (such as, for example, the ability to bind antigens or complement); and the ability to act as an antigen in a vaccine composition to raise an immune response against such protein or another material or entity which is cross-reactive with such protein.

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4.10.19 IDENTIFICATION OF POLYMORPHISMS

The demonstration of polymorphisms makes possible the identification of such polymorphisms in human subjects and the pharmacogenetic use of this information for diagnosis and treatment. Such polymorphisms may be associated with, e.g., differential predisposition or susceptibility to various disease states (such as disorders involving inflammation or immune response) or a differential response to drug administration, and this genetic information can be used to tailor preventive or therapeutic treatment appropriately. For example, the existence of a polymorphism associated with a predisposition to inflammation or autoimmune disease makes possible the diagnosis of this condition in humans by identifying the presence of the polymorphism.

Polymorphisms can be identified in a variety of ways known in the art which all generally involve obtaining a sample from a patient, analyzing DNA from the sample, optionally involving isolation or amplification of the DNA, and identifying the presence of the polymorphism in the DNA. For example, PCR may be used to amplify an appropriate fragment of genomic DNA which may then be sequenced. Alternatively, the DNA may be subjected to allele-specific oligonucleotide hybridization (in which

appropriate oligonucleotides are hybridized to the DNA under conditions permitting detection of a single base mismatch) or to a single nucleotide extension assay (in which an oligonucleotide that hybridizes immediately adjacent to the position of the polymorphism is extended with one or more labeled nucleotides). In addition, traditional restriction fragment length polymorphism analysis (using restriction enzymes that provide differential digestion of the genomic DNA depending on the presence or absence of the polymorphism) may be performed. Arrays with nucleotide sequences of the present invention can be used to detect polymorphisms. The array can comprise modified nucleotide sequences of the present invention. In the alternative, any one of the nucleotide sequences of the present invention can be placed on the array to detect changes from those sequences.

Alternatively a polymorphism resulting in a change in the amino acid sequence could also be detected by detecting a corresponding change in amino acid sequence of the protein, e.g., by an antibody specific to the variant sequence.

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4.10.20 ARTHRITIS AND INFLAMMATION

The immunosuppressive effects of the compositions of the invention against rheumatoid arthritis is determined in an experimental animal model system. The experimental model system is adjuvant induced arthritis in rats, and the protocol is described by J. Holoshitz, et at., 1983, Science, 219:56, or by B. Waksman et al., 1963, Int. Arch. Allergy Appl. Immunol., 23:129. Induction of the disease can be caused by a single injection, generally intradermally, of a suspension of killed Mycobacterium tuberculosis in complete Freund's adjuvant (CFA). The route of injection can vary, but rats may be injected at the base of the tail with an adjuvant mixture. The polypeptide is administered in phosphate buffered solution (PBS) at a dose of about 1-5 mg/kg. The control consists of administering PBS only.

The procedure for testing the effects of the test compound would consist of intradermally injecting killed Mycobacterium tuberculosis in CFA followed by immediately administering the test compound and subsequent treatment every other day until day 24. At 14, 15, 18, 20, 22, and 24 days after injection of Mycobacterium CFA, an overall arthritis score may be obtained as described by J. Holoskitz above. An analysis of

the data would reveal that the test compound would have a dramatic affect on the swelling of the joints as measured by a decrease of the arthritis score.

5 4.11 THERAPEUTIC METHODS

The compositions (including polypeptide fragments, analogs, variants and antibodies or other binding partners or modulators including antisense polynucleotides) of the invention have numerous applications in a variety of therapeutic methods. Examples of therapeutic applications include, but are not limited to, those exemplified herein.

4.11.1 EXAMPLE

One embodiment of the invention is the administration of an effective amount of the polypeptides or other composition of the invention to individuals affected by a disease or disorder that can be modulated by regulating the peptides of the invention. While the mode of administration is not particularly important, parenteral administration is preferred. An exemplary mode of administration is to deliver an intravenous bolus. The dosage of the polypeptides or other composition of the invention will normally be determined by the prescribing physician. It is to be expected that the dosage will vary according to the age, weight, condition and response of the individual patient. Typically, the amount of polypeptide administered per dose will be in the range of about 0.01µg/kg to 100 mg/kg of body weight, with the preferred dose being about 0.1µg/kg to 10 mg/kg of patient body weight. For parenteral administration, polypeptides of the invention will be formulated in an injectable form combined with a pharmaceutically acceptable parenteral vehicle. Such vehicles are well known in the art and examples include water, saline, Ringer's solution, dextrose solution, and solutions consisting of small amounts of the human serum albumin. The vehicle may contain minor amounts of additives that maintain the isotonicity and stability of the polypeptide or other active ingredient. The preparation of such solutions is within the skill of the art.

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4.12 PHARMACEUTICAL FORMULATIONS AND ROUTES OF ADMINISTRATION

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A protein or other composition of the present invention (from whatever source derived, including without limitation from recombinant and non-recombinant sources and including antibodies and other binding partners of the polypeptides of the invention) may be administered to a patient in need, by itself, or in pharmaceutical compositions where it is mixed with suitable carriers or excipient(s) at doses to treat or ameliorate a variety of disorders. Such a composition may optionally contain (in addition to protein or other active ingredient and a carrier) diluents, fillers, salts, buffers, stabilizers, solubilizers, and other materials well known in the art. The term "pharmaceutically acceptable" means a non-toxic material that does not interfere with the effectiveness of the biological activity of the active ingredient(s). The characteristics of the carrier will depend on the route of administration. The pharmaceutical composition of the invention may also contain cytokines, lymphokines, or other hematopoietic factors such as M-CSF, GM-CSF, TNF, IL-1, IL-2, IL-3, IL-4, IL-5, IL-6, IL-7, IL-8, IL-9, IL-10, IL-11, IL-12, IL-13, IL-14, IL-15, IFN, TNF0, TNF1, TNF2, G-CSF, Meg-CSF, thrombopoietin, stem cell factor, and erythropoietin. In further compositions, proteins of the invention may be combined with other agents beneficial to the treatment of the disease or disorder in question. These agents include various growth factors such as epidermal growth factor (EGF), platelet-derived growth factor (PDGF), transforming growth factors (TGF-\alpha and TGF-\beta), insulin-like growth factor (IGF), as well as cytokines described herein.

The pharmaceutical composition may further contain other agents which either enhance the activity of the protein or other active ingredient or complement its activity or use in treatment. Such additional factors and/or agents may be included in the pharmaceutical composition to produce a synergistic effect with protein or other active ingredient of the invention, or to minimize side effects. Conversely, protein or other active ingredient of the present invention may be included in formulations of the particular clotting factor, cytokine, lymphokine, other hematopoietic factor, thrombolytic or anti-thrombotic factor, or anti-inflammatory agent to minimize side effects of the clotting factor, cytokine, lymphokine, other hematopoietic factor, thrombolytic or anti-thrombotic factor, or anti-inflammatory agent (such as IL-1Ra, IL-1 Hy1, IL-1 Hy2,

anti-TNF, corticosteroids, immunosuppressive agents). A protein of the present invention may be active in multimers (e.g., heterodimers or homodimers) or complexes with itself or other proteins. As a result, pharmaceutical compositions of the invention may comprise a protein of the invention in such multimeric or complexed form.

As an alternative to being included in a pharmaceutical composition of the invention including a first protein, a second protein or a therapeutic agent may be concurrently administered with the first protein (e.g., at the same time, or at differing times provided that therapeutic concentrations of the combination of agents is achieved at the treatment site). Techniques for formulation and administration of the compounds of the instant application may be found in "Remington's Pharmaceutical Sciences," Mack Publishing Co., Easton, PA, latest edition. A therapeutically effective dose further refers to that amount of the compound sufficient to result in amelioration of symptoms, e.g., treatment, healing, prevention or amelioration of the relevant medical condition, or an increase in rate of treatment, healing, prevention or amelioration of such conditions. When applied to an individual active ingredient, administered alone, a therapeutically effective dose refers to that ingredient alone. When applied to a combination, a therapeutically effective dose refers to combined amounts of the active ingredients that result in the therapeutic effect, whether administered in combination, serially or simultaneously.

In practicing the method of treatment or use of the present invention, a therapeutically effective amount of protein or other active ingredient of the present invention is administered to a mammal having a condition to be treated. Protein or other active ingredient of the present invention may be administered in accordance with the method of the invention either alone or in combination with other therapies such as treatments employing cytokines, lymphokines or other hematopoietic factors. When coadministered with one or more cytokines, lymphokines or other hematopoietic factors, protein or other active ingredient of the present invention may be administered either simultaneously with the cytokine(s), lymphokine(s), other hematopoietic factor(s), thrombolytic or anti-thrombotic factors, or sequentially. If administered sequentially, the attending physician will decide on the appropriate sequence of administering protein or

other active ingredient of the present invention in combination with cytokine(s), lymphokine(s), other hematopoietic factor(s), thrombolytic or anti-thrombotic factors.

4.12.1 ROUTES OF ADMINISTRATION

Suitable routes of administration may, for example, include oral, rectal, transmucosal, or intestinal administration; parenteral delivery, including intramuscular, subcutaneous, intramedullary injections, as well as intrathecal, direct intraventricular, intravenous, intraperitoneal, intranasal, or intraocular injections. Administration of protein or other active ingredient of the present invention used in the pharmaceutical composition or to practice the method of the present invention can be carried out in a variety of conventional ways, such as oral ingestion, inhalation, topical application or cutaneous, subcutaneous, intraperitoneal, parenteral or intravenous injection. Intravenous administration to the patient is preferred.

Alternately, one may administer the compound in a local rather than systemic manner, for example, via injection of the compound directly into a arthritic joints or in fibrotic tissue, often in a depot or sustained release formulation. In order to prevent the scarring process frequently occurring as complication of glaucoma surgery, the compounds may be administered topically, for example, as eye drops. Furthermore, one may administer the drug in a targeted drug delivery system, for example, in a liposome coated with a specific antibody, targeting, for example, arthritic or fibrotic tissue. The liposomes will be targeted to and taken up selectively by the afflicted tissue.

The polypeptides of the invention are administered by any route that delivers an effective dosage to the desired site of action. The determination of a suitable route of administration and an effective dosage for a particular indication is within the level of skill in the art. Preferably for wound treatment, one administers the therapeutic compound directly to the site. Suitable dosage ranges for the polypeptides of the invention can be extrapolated from these dosages or from similar studies in appropriate animal models. Dosages can then be adjusted as necessary by the clinician to provide maximal therapeutic benefit.

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4.12.2 COMPOSITIONS/FORMULATIONS

Pharmaceutical compositions for use in accordance with the present invention thus may be formulated in a conventional manner using one or more physiologically acceptable carriers comprising excipients and auxiliaries which facilitate processing of the active compounds into preparations which can be used pharmaceutically. These pharmaceutical compositions may be manufactured in a manner that is itself known, e.g., by means of conventional mixing, dissolving, granulating, dragee-making, levigating, emulsifying, encapsulating, entrapping or lyophilizing processes. Proper formulation is dependent upon the route of administration chosen. When a therapeutically effective amount of protein or other active ingredient of the present invention is administered orally, protein or other active ingredient of the present invention will be in the form of a tablet, capsule, powder, solution or elixir. When administered in tablet form, the pharmaceutical composition of the invention may additionally contain a solid carrier such as a gelatin or an adjuvant. The tablet, capsule, and powder contain from about 5 to 95% protein or other active ingredient of the present invention, and preferably from about 25 to 90% protein or other active ingredient of the present invention. When administered in liquid form, a liquid carrier such as water, petroleum, oils of animal or plant origin such as peanut oil, mineral oil, soybean oil, or sesame oil, or synthetic oils may be added. The liquid form of the pharmaceutical composition may further contain physiological saline solution, dextrose or other saccharide solution, or glycols such as ethylene glycol, propylene glycol or polyethylene glycol. When administered in liquid form, the pharmaceutical composition contains from about 0.5 to 90% by weight of protein or other active ingredient of the present invention, and preferably from about 1 to 50% protein or other active ingredient of the present invention.

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When a therapeutically effective amount of protein or other active ingredient of the present invention is administered by intravenous, cutaneous or subcutaneous injection, protein or other active ingredient of the present invention will be in the form of a pyrogen-free, parenterally acceptable aqueous solution. The preparation of such parenterally acceptable protein or other active ingredient solutions, having due regard to pH, isotonicity, stability, and the like, is within the skill in the art. A preferred 30 pharmaceutical composition for intravenous, cutaneous, or subcutaneous injection should contain, in addition to protein or other active ingredient of the present invention, an

isotonic vehicle such as Sodium Chloride Injection, Ringer's Injection, Dextrose
Injection, Dextrose and Sodium Chloride Injection, Lactated Ringer's Injection, or other
vehicle as known in the art. The pharmaceutical composition of the present invention
may also contain stabilizers, preservatives, buffers, antioxidants, or other additives
known to those of skill in the art. For injection, the agents of the invention may be
formulated in aqueous solutions, preferably in physiologically compatible buffers such as
Hanks's solution, Ringer's solution, or physiological saline buffer. For transmucosal
administration, penetrants appropriate to the barrier to be permeated are used in the
formulation. Such penetrants are generally known in the art.

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For oral administration, the compounds can be formulated readily by combining the active compounds with pharmaceutically acceptable carriers well known in the art. Such carriers enable the compounds of the invention to be formulated as tablets, pills, dragees, capsules, liquids, gels, syrups, slurries, suspensions and the like, for oral ingestion by a patient to be treated. Pharmaceutical preparations for oral use can be obtained from a solid excipient, optionally grinding a resulting mixture, and processing the mixture of granules, after adding suitable auxiliaries, if desired, to obtain tablets or dragee cores. Suitable excipients are, in particular, fillers such as sugars, including lactose, sucrose, mannitol, or sorbitol; cellulose preparations such as, for example, maize starch, wheat starch, rice starch, potato starch, gelatin, gum tragacanth, methyl cellulose, hydroxypropylmethyl-cellulose, sodium carboxymethylcellulose, and/or polyvinylpyrrolidone (PVP). If desired, disintegrating agents may be added, such as the cross-linked polyvinyl pyrrolidone, agar, or alginic acid or a salt thereof such as sodium alginate. Dragee cores are provided with suitable coatings. For this purpose, concentrated sugar solutions may be used, which may optionally contain gum arabic, talc, polyvinyl pyrrolidone, carbopol gel, polyethylene glycol, and/or titanium dioxide, lacquer solutions, and suitable organic solvents or solvent mixtures. Dyestuffs or pigments may be added to the tablets or dragee coatings for identification or to characterize different combinations of active compound doses.

Pharmaceutical preparations which can be used orally include push-fit capsules made of gelatin, as well as soft, sealed capsules made of gelatin and a plasticizer, such as glycerol or sorbitol. The push-fit capsules can contain the active ingredients in admixture

with filler such as lactose, binders such as starches, and/or lubricants such as talc or magnesium stearate and, optionally, stabilizers. In soft capsules, the active compounds may be dissolved or suspended in suitable liquids, such as fatty oils, liquid paraffin, or liquid polyethylene glycols. In addition, stabilizers may be added. All formulations for oral administration should be in dosages suitable for such administration. For buccal administration, the compositions may take the form of tablets or lozenges formulated in conventional manner.

For administration by inhalation, the compounds for use according to the present invention are conveniently delivered in the form of an aerosol spray presentation from pressurized packs or a nebuliser, with the use of a suitable propellant, e.g., dichlorodifluoromethane, trichlorofluoromethane, dichlorotetrafluoroethane, carbon dioxide or other suitable gas. In the case of a pressurized aerosol the dosage unit may be determined by providing a valve to deliver a metered amount. Capsules and cartridges of, e.g., gelatin for use in an inhaler or insufflator may be formulated containing a powder mix of the compound and a suitable powder base such as lactose or starch. The compounds may be formulated for parenteral administration by injection, e.g., by bolus injection or continuous infusion. Formulations for injection may be presented in unit dosage form, e.g., in ampules or in multi-dose containers, with an added preservative. The compositions may take such forms as suspensions, solutions or emulsions in oily or aqueous vehicles, and may contain formulatory agents such as suspending, stabilizing and/or dispersing agents.

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Pharmaceutical formulations for parenteral administration include aqueous solutions of the active compounds in water-soluble form. Additionally, suspensions of the active compounds may be prepared as appropriate oily injection suspensions. Suitable lipophilic solvents or vehicles include fatty oils such as sesame oil, or synthetic fatty acid esters, such as ethyl oleate or triglycerides, or liposomes. Aqueous injection suspensions may contain substances which increase the viscosity of the suspension, such as sodium carboxymethyl cellulose, sorbitol, or dextran. Optionally, the suspension may also contain suitable stabilizers or agents which increase the solubility of the compounds to allow for the preparation of highly concentrated solutions. Alternatively, the active

ingredient may be in powder form for constitution with a suitable vehicle, e.g., sterile pyrogen-free water, before use.

The compounds may also be formulated in rectal compositions such as suppositories or retention enemas, e.g., containing conventional suppository bases such as 5 - cocoa butter-or-other glycerides.—In addition to the formulations described previously, the compounds may also be formulated as a depot preparation. Such long acting formulations may be administered by implantation (for example subcutaneously or intramuscularly) or by intramuscular injection. Thus, for example, the compounds may be formulated with suitable polymeric or hydrophobic materials (for example as an emulsion in an acceptable oil) or ion exchange resins, or as sparingly soluble derivatives, for example, as a sparingly soluble salt.

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A pharmaceutical carrier for the hydrophobic compounds of the invention is a cosolvent system comprising benzyl alcohol, a nonpolar surfactant, a water-miscible organic polymer, and an aqueous phase. The co-solvent system may be the VPD 15 co-solvent system. VPD is a solution of 3% w/v benzyl alcohol, 8% w/v of the nonpolar surfactant polysorbate 80, and 65% w/v polyethylene glycol 300, made up to volume in absolute ethanol. The VPD co-solvent system (VPD:5W) consists of VPD diluted 1:1 with a 5% dextrose in water solution. This co-solvent system dissolves hydrophobic compounds well, and itself produces low toxicity upon systemic administration. 20 Naturally, the proportions of a co-solvent system may be varied considerably without destroying its solubility and toxicity characteristics. Furthermore, the identity of the co-solvent components may be varied: for example, other low-toxicity nonpolar surfactants may be used instead of polysorbate 80; the fraction size of polyethylene glycol may be varied; other biocompatible polymers may replace polyethylene glycol, 25 e.g. polyvinyl pyrrolidone; and other sugars or polysaccharides may substitute for dextrose. Alternatively, other delivery systems for hydrophobic pharmaceutical compounds may be employed. Liposomes and emulsions are well known examples of delivery vehicles or carriers for hydrophobic drugs. Certain organic solvents such as dimethylsulfoxide also may be employed, although usually at the cost of greater toxicity. 30 Additionally, the compounds may be delivered using a sustained-release system, such as semipermeable matrices of solid hydrophobic polymers containing the therapeutic agent.

Various types of sustained-release materials have been established and are well known by those skilled in the art. Sustained-release capsules may, depending on their chemical nature, release the compounds for a few weeks up to over 100 days. Depending on the chemical nature and the biological stability of the therapeutic reagent, additional strategies for protein or other active ingredient stabilization may be employed.

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The pharmaceutical compositions also may comprise suitable solid or gel phase carriers or excipients. Examples of such carriers or excipients include but are not limited to calcium carbonate, calcium phosphate, various sugars, starches, cellulose derivatives, gelatin, and polymers such as polyethylene glycols. Many of the active ingredients of the invention may be provided as salts with pharmaceutically compatible counter ions. Such pharmaceutically acceptable base addition salts are those salts which retain the biological effectiveness and properties of the free acids and which are obtained by reaction with inorganic or organic bases such as sodium hydroxide, magnesium hydroxide, ammonia, trialkylamine, dialkylamine, monoalkylamine, dibasic amino acids, sodium acetate, potassium benzoate, triethanol amine and the like.

The pharmaceutical composition of the invention may be in the form of a complex of the protein(s) or other active ingredient(s) of present invention along with protein or peptide antigens. The protein and/or peptide antigen will deliver a stimulatory signal to both B and T lymphocytes. B lymphocytes will respond to antigen through their surface immunoglobulin receptor. T lymphocytes will respond to antigen through the T cell receptor (TCR) following presentation of the antigen by MHC proteins. MHC and structurally related proteins including those encoded by class I and class II MHC genes on host cells will serve to present the peptide antigen(s) to T lymphocytes. The antigen components could also be supplied as purified MHC-peptide complexes alone or with co-stimulatory molecules that can directly signal T cells. Alternatively antibodies able to bind surface immunoglobulin and other molecules on B cells as well as antibodies able to bind the TCR and other molecules on T cells can be combined with the pharmaceutical composition of the invention.

The pharmaceutical composition of the invention may be in the form of a liposome in which protein of the present invention is combined, in addition to other pharmaceutically acceptable carriers, with amphipathic agents such as lipids which exist

in aggregated form as micelles, insoluble monolayers, liquid crystals, or lamellar layers in aqueous solution. Suitable lipids for liposomal formulation include, without limitation, monoglycerides, diglycerides, sulfatides, lysolecithins, phospholipids, saponin, bile acids, and the like. Preparation of such liposomal formulations is within the level of skill in the art, as disclosed, for example, in U.S. Patent Nos. 4,235,871; 4,501,728; 4,837,028; and 4,737,323, all of which are incorporated herein by reference.

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The amount of protein or other active ingredient of the present invention in the pharmaceutical composition of the present invention will depend upon the nature and severity of the condition being treated, and on the nature of prior treatments which the patient has undergone. Ultimately, the attending physician will decide the amount of protein or other active ingredient of the present invention with which to treat each individual patient. Initially, the attending physician will administer low doses of protein or other active ingredient of the present invention and observe the patient's response. Larger doses of protein or other active ingredient of the present invention may be administered until the optimal therapeutic effect is obtained for the patient, and at that point the dosage is not increased further. It is contemplated that the various pharmaceutical compositions used to practice the method of the present invention should contain about 0.01 µg to about 100 mg (preferably about 0.1 µg to about 10 mg, more preferably about 0.1 µg to about 1 mg) of protein or other active ingredient of the present invention per kg body weight. For compositions of the present invention which are useful for bone, cartilage, tendon or ligament regeneration, the therapeutic method includes administering the composition topically, systematically, or locally as an implant or device. When administered, the therapeutic composition for use in this invention is, of course, in a pyrogen-free, physiologically acceptable form. Further, the composition may desirably be encapsulated or injected in a viscous form for delivery to the site of bone, cartilage or tissue damage. Topical administration may be suitable for wound healing and tissue repair. Therapeutically useful agents other than a protein or other active ingredient of the invention which may also optionally be included in the composition as described above, may alternatively or additionally, be administered simultaneously or sequentially with the composition in the methods of the invention. Preferably for bone and/or cartilage formation, the composition would include a matrix capable of delivering

the protein-containing or other active ingredient-containing composition to the site of bone and/or cartilage damage, providing a structure for the developing bone and cartilage and optimally capable of being resorbed into the body. Such matrices may be formed of materials presently in use for other implanted medical applications.

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The choice of matrix material is based on biocompatibility, biodegradability, mechanical properties, cosmetic appearance and interface properties. The particular application of the compositions will define the appropriate formulation. Potential matrices for the compositions may be biodegradable and chemically defined calcium sulfate, tricalcium phosphate, hydroxyapatite, polylactic acid, polyglycolic acid and polyanhydrides. Other potential materials are biodegradable and biologically well-defined, such as bone or dermal collagen. Further matrices are comprised of pure proteins or extracellular matrix components. Other potential matrices are nonbiodegradable and chemically defined, such as sintered hydroxyapatite, bioglass. aluminates, or other ceramics. Matrices may be comprised of combinations of any of the above mentioned types of material, such as polylactic acid and hydroxyapatite or collagen and tricalcium phosphate. The bioceramics may be altered in composition, such as in calcium-aluminate-phosphate and processing to alter pore size, particle size, particle shape, and biodegradability. Presently preferred is a 50:50 (mole weight) copolymer of lactic acid and glycolic acid in the form of porous particles having diameters ranging from 150 to 800 microns. In some applications, it will be useful to utilize a sequestering agent, such as carboxymethyl cellulose or autologous blood clot, to prevent the protein compositions from disassociating from the matrix.

A preferred family of sequestering agents is cellulosic materials such as alkylcelluloses (including hydroxyalkylcelluloses), including methylcellulose, ethylcellulose, hydroxypropylcellulose, hydroxypropylcellulose, hydroxypropyl-methylcellulose, and carboxymethylcellulose, the most preferred being cationic salts of carboxymethylcellulose (CMC). Other preferred sequestering agents include hyaluronic acid, sodium alginate, poly(ethylene glycol), polyoxyethylene oxide, carboxyvinyl polymer and poly(vinyl alcohol). The amount of sequestering agent useful herein is 0.5-20 wt %, preferably 1-10 wt % based on total formulation weight, which represents the amount necessary to prevent desorption of the protein from the polymer

matrix and to provide appropriate handling of the composition, yet not so much that the progenitor cells are prevented from infiltrating the matrix, thereby providing the protein the opportunity to assist the osteogenic activity of the progenitor cells. In further compositions, proteins or other active ingredients of the invention may be combined with other agents beneficial to the treatment of the bone and/or cartilage defect, wound, or tissue in question. These agents include various growth factors such as epidermal growth factor (EGF), platelet derived growth factor (PDGF), transforming growth factors (TGF-α and TGF-β), and insulin-like growth factor (IGF).

The therapeutic compositions are also presently valuable for veterinary applications. Particularly domestic animals and thoroughbred horses, in addition to humans, are desired patients for such treatment with proteins or other active ingredients of the present invention. The dosage regimen of a protein-containing pharmaceutical composition to be used in tissue regeneration will be determined by the attending physician considering various factors which modify the action of the proteins, e.g., amount of tissue weight desired to be formed, the site of damage, the condition of the damaged tissue, the size of a wound, type of damaged tissue (e.g., bone), the patient's age, sex, and diet, the severity of any infection, time of administration and other clinical factors. The dosage may vary with the type of matrix used in the reconstitution and with inclusion of other proteins in the pharmaceutical composition. For example, the addition of other known growth factors, such as IGF I (insulin like growth factor I), to the final composition, may also effect the dosage. Progress can be monitored by periodic assessment of tissue/bone growth and/or repair, for example, X-rays, histomorphometric determinations and tetracycline labeling.

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Polynucleotides of the present invention can also be used for gene therapy. Such polynucleotides can be introduced either in vivo or ex vivo into cells for expression in a mammalian subject. Polynucleotides of the invention may also be administered by other known methods for introduction of nucleic acid into a cell or organism (including, without limitation, in the form of viral vectors or naked DNA). Cells may also be cultured ex vivo in the presence of proteins of the present invention in order to proliferate or to produce a desired effect on or activity in such cells. Treated cells can then be introduced in vivo for therapeutic purposes.

4.12.3 EFFECTIVE DOSAGE

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Pharmaceutical compositions suitable for use in the present invention include compositions wherein the active ingredients are contained in an effective amount to achieve its intended purpose. More specifically, a therapeutically effective amount means an amount effective to prevent development of or to alleviate the existing symptoms of the subject being treated. Determination of the effective amount is well within the capability of those skilled in the art, especially in light of the detailed disclosure provided herein. For any compound used in the method of the invention, the therapeutically effective dose can be estimated initially from appropriate in vitro assays. For example, a dose can be formulated in animal models to achieve a circulating concentration range that can be used to more accurately determine useful doses in humans. For example, a dose can be formulated in animal models to achieve a circulating concentration range that includes the IC₅₀ as determined in cell culture (i.e., the concentration of the test compound which achieves a half-maximal inhibition of the protein's biological activity). Such information can be used to more accurately determine useful doses in humans.

A therapeutically effective dose refers to that amount of the compound that results in amelioration of symptoms or a prolongation of survival in a patient. Toxicity and 20 therapeutic efficacy of such compounds can be determined by standard pharmaceutical procedures in cell cultures or experimental animals, e.g., for determining the LD₅₀ (the dose lethal to 50% of the population) and the ED₅₀ (the dose therapeutically effective in 50% of the population). The dose ratio between toxic and therapeutic effects is the therapeutic index and it can be expressed as the ratio between LD₅₀ and ED₅₀. 25 Compounds which exhibit high therapeutic indices are preferred. The data obtained from these cell culture assays and animal studies can be used in formulating a range of dosage for use in human. The dosage of such compounds lies preferably within a range of circulating concentrations that include the ED₅₀ with little or no toxicity. The dosage may vary within this range depending upon the dosage form employed and the route of 30 administration utilized. The exact formulation, route of administration and dosage can be chosen by the individual physician in view of the patient's condition. See, e.g., Fingl et

al., 1975, in "The Pharmacological Basis of Therapeutics", Ch. 1 p.1. Dosage amount and interval may be adjusted individually to provide plasma levels of the active moiety which are sufficient to maintain the desired effects, or minimal effective concentration (MEC). The MEC will vary for each compound but can be estimated from *in vitro* data. Dosages necessary to achieve the MEC will depend on individual characteristics and route of administration. However, HPLC assays or bioassays can be used to determine plasma concentrations.

Dosage intervals can also be determined using MEC value. Compounds should be administered using a regimen which maintains plasma levels above the MEC for 10-90% of the time, preferably between 30-90% and most preferably between 50-90%. In cases of local administration or selective uptake, the effective local concentration of the drug may not be related to plasma concentration.

An exemplary dosage regimen for polypeptides or other compositions of the invention will be in the range of about 0.01 μ g/kg to 100 mg/kg of body weight daily, with the preferred dose being about 0.1 μ g/kg to 25 mg/kg of patient body weight daily, varying in adults and children. Dosing may be once daily, or equivalent doses may be delivered at longer or shorter intervals.

The amount of composition administered will, of course, be dependent on the subject being treated, on the subject's age and weight, the severity of the affliction, the manner of administration and the judgment of the prescribing physician.

4.12.4 PACKAGING

The compositions may, if desired, be presented in a pack or dispenser device which may contain one or more unit dosage forms containing the active ingredient. The pack may, for example, comprise metal or plastic foil, such as a blister pack. The pack or dispenser device may be accompanied by instructions for administration. Compositions comprising a compound of the invention formulated in a compatible pharmaceutical carrier may also be prepared, placed in an appropriate container, and labeled for treatment of an indicated condition.

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4.13 ANTIBODIES

Also included in the invention are antibodies to proteins, or fragments of proteins of the invention. The term "antibody" as used herein refers to immunoglobulin molecules and immunologically active portions of immunoglobulin (Ig) molecules, i.e., molecules that contain an antigen-binding site that specifically binds (immunoreacts with) an antigen. Such antibodies include, but are not limited to, polyclonal, monoclonal, chimeric, single chain, F_{ab} , $F_{ab'}$ and $F_{(ab')2}$ fragments, and an F_{ab} expression library. In general, an antibody molecule obtained from humans relates to any of the classes IgG, IgM, IgA, IgE and IgD, which differ from one another by the nature of the heavy chain present in the molecule. Certain classes have subclasses as well, such as IgG₁, IgG₂, and others. Furthermore, in humans, the light chain may be a kappa chain or a lambda chain. Reference herein to antibodies includes a reference to all such classes, subclasses and types of human antibody species.

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An isolated related protein of the invention may be intended to serve as an antigen, or a portion or fragment thereof, and additionally can be used as an immunogen to generate antibodies that immunospecifically bind the antigen, using standard techniques for polyclonal and monoclonal antibody preparation. The full-length protein can be used or, alternatively, the invention provides antigenic peptide fragments of the antigen for use as immunogens. An antigenic peptide fragment comprises at least 6 amino acid residues of the amino acid sequence of the full length protein, such as an amino acid sequence shown in SEQ ID NO: 1-438, and encompasses an epitope thereof such that an antibody raised against the peptide forms a specific immune complex with the full length protein or with any fragment that contains the epitope. Preferably, the antigenic peptide comprises at least 10 amino acid residues, or at least 15 amino acid residues, or at least 20 amino acid residues, or at least 30 amino acid residues. Preferred epitopes encompassed by the antigenic peptide are regions of the protein that are located on its surface; commonly these are hydrophilic regions.

In certain embodiments of the invention, at least one epitope encompassed by the antigenic peptide is a region of alpha-2-macroglobulin-like protein that is located on the surface of the protein, e.g., a hydrophilic region. A hydrophobicity analysis of the human related protein sequence will indicate which regions of a related protein are particularly hydrophilic and, therefore, are likely to encode surface residues useful for targeting

antibody production. As a means for targeting antibody production, hydropathy plots showing regions of hydrophilicity and hydrophobicity may be generated by any method well known in the art, including, for example, the Kyte Doolittle or the Hopp Woods methods, either with or without Fourier transformation. See, e.g., Hopp and Woods,

1981, Proc. Nat. Acad. Sci. USA 78: 3824-3828; Kyte and Doolittle 1982, J. Mol. Biol. 157: 105-142, each of which is incorporated herein by reference in its entirety. Antibodies that are specific for one or more domains within an antigenic protein, or derivatives, fragments, analogs or homologs thereof, are also provided herein.

A protein of the invention, or a derivative, fragment, analog, homolog or ortholog thereof, may be utilized as an immunogen in the generation of antibodies that immunospecifically bind these protein components.

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The term "specific for" indicates that the variable regions of the antibodies of the invention recognize and bind polypeptides of the invention exclusively (i.e., able to distinguish the polypeptide of the invention from other similar polypeptides despite sequence identity, homology, or similarity found in the family of polypeptides), but may also interact with other proteins (for example, S. aureus protein A or other antibodies in ELISA techniques) through interactions with sequences outside the variable region of the antibodies, and in particular, in the constant region of the molecule. Screening assays to determine binding specificity of an antibody of the invention are well known and routinely practiced in the art. For a comprehensive discussion of such assays, see Harlow et al. (Eds), Antibodies A Laboratory Manual; Cold Spring Harbor Laboratory; Cold Spring Harbor, NY (1988), Chapter 6. Antibodies that recognize and bind fragments of the polypeptides of the invention are also contemplated, provided that the antibodies are first and foremost specific for, as defined above, full-length polypeptides of the invention. As with antibodies that are specific for full length polypeptides of the invention, antibodies of the invention that recognize fragments are those which can distinguish polypeptides from the same family of polypeptides despite inherent sequence identity, homology, or similarity found in the family of proteins.

Antibodies of the invention are useful for, for example, therapeutic purposes (by modulating activity of a polypeptide of the invention), diagnostic purposes to detect or quantitate a polypeptide of the invention, as well as purification of a polypeptide of the

invention. Kits comprising an antibody of the invention for any of the purposes described herein are also comprehended. In general, a kit of the invention also includes a control antigen for which the antibody is immunospecific. The invention further provides a hybridoma that produces an antibody according to the invention. Antibodies of the invention are useful for detection and/or purification of the polypeptides of the invention.

Monoclonal antibodies binding to the protein of the invention may be useful diagnostic agents for the immunodetection of the protein. Neutralizing monoclonal antibodies binding to the protein may also be useful therapeutics for both conditions associated with the protein and also in the treatment of some forms of cancer where abnormal expression of the protein is involved. In the case of cancerous cells or leukemic cells, neutralizing monoclonal antibodies against the protein may be useful in detecting and preventing the metastatic spread of the cancerous cells, which may be mediated by the protein.

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The labeled antibodies of the present invention can be used for *in vitro*, *in vivo*, and *in situ* assays to identify cells or tissues in which a fragment of the polypeptide of interest is expressed. The antibodies may also be used directly in therapies or other diagnostics. The present invention further provides the above-described antibodies immobilized on a solid support. Examples of such solid supports include plastics such as polycarbonate, complex carbohydrates such as agarose and Sepharose®, acrylic resins and such as polyacrylamide and latex beads. Techniques for coupling antibodies to such solid supports are well known in the art (Weir, D.M. et al., "Handbook of Experimental Immunology" 4th Ed., Blackwell Scientific Publications, Oxford, England, Chapter 10 (1986); Jacoby, W.D. et al., Meth. Enzym. 34 Academic Press, N.Y. (1974)). The immobilized antibodies of the present invention can be used for *in vitro*, *in vivo*, and *in situ* assays as well as for immuno-affinity purification of the proteins of the present invention.

Various procedures known within the art may be used for the production of polyclonal or monoclonal antibodies directed against a protein of the invention, or against derivatives, fragments, analogs homologs or orthologs thereof (see, for example, Antibodies: A Laboratory Manual, Harlow E, and Lane D, 1988, Cold Spring Harbor

Laboratory Press, Cold Spring Harbor, NY, incorporated herein by reference). Some of these antibodies are discussed below.

4.13.1 POLYCLONAL ANTIBODIES

5 For the production of polyclonal antibodies, various suitable host animals (e.g., rabbit, goat, mouse or other mammal) may be immunized by one or more injections with the native protein, a synthetic variant thereof, or a derivative of the foregoing. An appropriate immunogenic preparation can contain, for example, the naturally occurring immunogenic protein, a chemically synthesized polypeptide representing the immunogenic protein, or a recombinantly expressed immunogenic protein. Furthermore, 10 the protein may be conjugated to a second protein known to be immunogenic in the mammal being immunized. Examples of such immunogenic proteins include but are not limited to keyhole limpet hemocyanin, serum albumin, bovine thyroglobulin, and soybean trypsin inhibitor. The preparation can further include an adjuvant. Various adjuvants used to increase the immunological response include, but are not limited to, 15 Freund's (complete and incomplete), mineral gels (e.g., aluminum hydroxide), surfaceactive substances (e.g., lysolecithin, pluronic polyols, polyanions, peptides, oil emulsions, dinitrophenol, etc.), adjuvants usable in humans such as Bacille Calmette-Guerin and Corynebacterium parvum, or similar immunostimulatory agents. Additional examples of 20 adjuvants that can be employed include MPL-TDM adjuvant (monophosphoryl Lipid A, synthetic trehalose dicorynomycolate).

The polyclonal antibody molecules directed against the immunogenic protein can be isolated from the mammal (e.g., from the blood) and further purified by well known techniques, such as affinity chromatography using protein A or protein G, which provide primarily the IgG fraction of immune serum. Subsequently, or alternatively, the specific antigen which is the target of the immunoglobulin sought, or an epitope thereof, may be immobilized on a column to purify the immune specific antibody by immunoaffinity chromatography. Purification of immunoglobulins is discussed, for example, by D. Wilkinson (The Scientist, published by The Scientist, Inc., Philadelphia PA, Vol. 14, No. 8 (April 17, 2000), pp. 25-28).

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4.13.2 MONOCLONAL ANTIBODIES

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The term "monoclonal antibody" (MAb) or "monoclonal antibody composition", as used herein, refers to a population of antibody molecules that contain only one molecular species of antibody molecule consisting of a unique light chain gene product and a unique heavy chain gene product. In particular, the complementarity determining regions (CDRs) of the monoclonal antibody are identical in all the molecules of the population. MAbs thus contain an antigen-binding site capable of immunoreacting with a particular epitope of the antigen characterized by a unique binding affinity for it.

Monoclonal antibodies can be prepared using hybridoma methods, such as those described by Kohler and Milstein, Nature, 256:495 (1975). In a hybridoma method, a mouse, hamster, or other appropriate host animal, is typically immunized with an immunizing agent to elicit lymphocytes that produce or are capable of producing antibodies that will specifically bind to the immunizing agent. Alternatively, the lymphocytes can be immunized in vitro.

The immunizing agent will typically include the protein antigen, a fragment thereof or a fusion protein thereof. Generally, either peripheral blood lymphocytes are used if cells of human origin are desired, or spleen cells or lymph node cells are used if non-human mammalian sources are desired. The lymphocytes are then fused with an immortalized cell line using a suitable fusing agent, such as polyethylene glycol, to form a hybridoma cell (Goding, Monoclonal Antibodies: Principles and Practice, Academic Press, (1986) pp. 59-103). Immortalized cell lines are usually transformed mammalian cells, particularly myeloma cells of rodent, bovine and human origin. Usually, rat or mouse myeloma cell lines are employed. The hybridoma cells can be cultured in a suitable culture medium that preferably contains one or more substances that inhibit the growth or survival of the unfused, immortalized cells. For example, if the parental cells lack the enzyme hypoxanthine guanine phosphoribosyl transferase (HGPRT or HPRT), the culture medium for the hybridomas typically will include hypoxanthine, aminopterin, and thymidine ("HAT medium"), which substances prevent the growth of HGPRT-deficient cells.

Preferred immortalized cell lines are those that fuse efficiently, support stable high level expression of antibody by the selected antibody-producing cells, and are

sensitive to a medium such as HAT medium. More preferred immortalized cell lines are murine myeloma lines, which can be obtained, for instance, from the Salk Institute Cell Distribution Center, San Diego, California and the American Type Culture Collection, Manassas, Virginia. Human myeloma and mouse-human heteromyeloma cell lines also have been described for the production of human monoclonal antibodies (Kozbor, <u>J. Immunol.</u>, <u>133</u>:3001 (1984); Brodeur et al., <u>Monoclonal Antibody Production Techniques and Applications</u>, Marcel Dekker, Inc., New York, (1987) pp. 51-63).

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The culture medium in which the hybridoma cells are cultured can then be assayed for the presence of monoclonal antibodies directed against the antigen. Preferably, the binding specificity of monoclonal antibodies produced by the hybridoma cells is determined by immunoprecipitation or by an in vitro binding assay, such as radioimmunoassay (RIA) or enzyme-linked immunoabsorbent assay (ELISA). Such techniques and assays are known in the art. The binding affinity of the monoclonal antibody can, for example, be determined by the Scatchard analysis of Munson and Pollard, Anal. Biochem., 107:220 (1980). Preferably, antibodies having a high degree of specificity and a high binding affinity for the target antigen are isolated.

After the desired hybridoma cells are identified, the clones can be subcloned by limiting dilution procedures and grown by standard methods. Suitable culture media for this purpose include, for example, Dulbecco's Modified Eagle's Medium and RPMI-1640 medium. Alternatively, the hybridoma cells can be grown in vivo as ascites in a mammal.

The monoclonal antibodies secreted by the subclones can be isolated or purified from the culture medium or ascites fluid by conventional immunoglobulin purification procedures such as, for example, protein A-Sepharose, hydroxylapatite chromatography, gel electrophoresis, dialysis, or affinity chromatography.

The monoclonal antibodies can also be made by recombinant DNA methods, such as those described in U.S. Patent No. 4,816,567. DNA encoding the monoclonal antibodies of the invention can be readily isolated and sequenced using conventional procedures (e.g., by using oligonucleotide probes that are capable of binding specifically to genes encoding the heavy and light chains of murine antibodies). The hybridoma cells of the invention serve as a preferred source of such DNA. Once isolated, the DNA can

be placed into expression vectors, which are then transfected into host cells such as simian COS cells, Chinese hamster ovary (CHO) cells, or myeloma cells that do not otherwise produce immunoglobulin protein, to obtain the synthesis of monoclonal antibodies in the recombinant host cells. The DNA also can be modified, for example, by substituting the coding sequence for human heavy and light chain constant domains in place of the homologous murine sequences (U.S. Patent No. 4,816,567; Morrison, Nature 368, 812-13 (1994)) or by covalently joining to the immunoglobulin coding sequence all or part of the coding sequence for a non-immunoglobulin polypeptide. Such a non-immunoglobulin polypeptide can be substituted for the constant domains of an antibody of the invention, or can be substituted for the variable domains of one antigen-combining site of an antibody of the invention to create a chimeric bivalent antibody.

4.13.3 HUMANIZED ANTIBODIES

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The antibodies directed against the protein antigens of the invention can further 15 comprise humanized antibodies or human antibodies. These antibodies are suitable for administration to humans without engendering an immune response by the human against the administered immunoglobulin. Humanized forms of antibodies are chimeric immunoglobulins, immunoglobulin chains or fragments thereof (such as Fv, Fab, Fab', F(ab')₂ or other antigen-binding subsequences of antibodies) that are principally 20 comprised of the sequence of a human immunoglobulin, and contain minimal sequence derived from a non-human immunoglobulin. Humanization can be performed following the method of Winter and co-workers (Jones et al., Nature, 321:522-525 (1986); Riechmann et al., Nature, 332:323-327 (1988); Verhoeyen et al., Science, 239:1534-1536 (1988)), by substituting rodent CDRs or CDR sequences for the corresponding sequences 25 of a human antibody. (See also U.S. Patent No. 5,225,539). In some instances, Fv framework residues of the human immunoglobulin are replaced by corresponding nonhuman residues. Humanized antibodies can also comprise residues that are found neither in the recipient antibody nor in the imported CDR or framework sequences. In general, the humanized antibody will comprise substantially all of at least one, and typically two, 30 variable domains, in which all or substantially all of the CDR regions correspond to those of a non-human immunoglobulin and all or substantially all of the framework regions are

those of a human immunoglobulin consensus sequence. The humanized antibody optimally also will comprise at least a portion of an immunoglobulin constant region (Fc), typically that of a human immunoglobulin (Jones et al., 1986; Riechmann et al., 1988; and Presta, Curr. Op. Struct. Biol., 2:593-596 (1992)).

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4.13.4 HUMAN ANTIBODIES

Fully human antibodies relate to antibody molecules in which essentially the entire sequences of both the light chain and the heavy chain, including the CDRs, arise from human genes. Such antibodies are termed "human antibodies", or "fully human antibodies" herein. Human monoclonal antibodies can be prepared by the trioma technique; the human B-cell hybridoma technique (see Kozbor, et al., 1983 Immunol Today 4: 72) and the EBV hybridoma technique to produce human monoclonal antibodies (see Cole, et al., 1985 In: MONOCLONAL ANTIBODIES AND CANCER THERAPY, Alan R. Liss, Inc., pp. 77-96). Human monoclonal antibodies may be utilized in the practice of the present invention and may be produced by using human hybridomas (see Cote, et al., 1983. Proc Natl Acad Sci USA 80: 2026-2030) or by transforming human B-cells with Epstein Barr Virus in vitro (see Cole, et al., 1985 In: MONOCLONAL ANTIBODIES AND CANCER THERAPY, Alan R. Liss, Inc., pp. 77-96).

In addition, human antibodies can also be produced using additional techniques, including phage display libraries (Hoogenboom and Winter, J. Mol. Biol., 227:381 (1991); Marks et al., J. Mol. Biol., 222:581 (1991)). Similarly, human antibodies can be made by introducing human immunoglobulin loci into transgenic animals, e.g., mice in which the endogenous immunoglobulin genes have been partially or completely inactivated. Upon challenge, human antibody production is observed, which closely resembles that seen in humans in all respects, including gene rearrangement, assembly, and antibody repertoire. This approach is described, for example, in U.S. Patent Nos. 5,545,807; 5,545,806; 5,569,825; 5,625,126; 5,633,425; 5,661,016, and in Marks et al. (Bio/Technology 10, 779-783 (1992)); Lonberg et al. (Nature 368 856-859 (1994)); Morrison (Nature 368, 812-13 (1994)); Fishwild et al, (Nature Biotechnology 14, 845-51 (1996)); Neuberger (Nature Biotechnology 14, 826 (1996)); and Lonberg and Huszar (Intern. Rev. Immunol. 13 65-93 (1995)).

Human antibodies may additionally be produced using transgenic nonhuman animals that are modified so as to produce fully human antibodies rather than the animal's endogenous antibodies in response to challenge by an antigen. (See PCT publication WO94/02602). The endogenous genes encoding the heavy and light immunoglobulin chains in the nonhuman host have been incapacitated, and active loci encoding human heavy and light chain immunoglobulins are inserted into the host's genome. The human genes are incorporated, for example, using yeast artificial chromosomes containing the requisite human DNA segments. An animal which provides all the desired modifications is then obtained as progeny by crossbreeding intermediate transgenic animals containing fewer than the full complement of the modifications. The preferred embodiment of such a nonhuman animal is a mouse, and is termed the Xenomouse[™] as disclosed in PCT publications WO 96/33735 and WO 96/34096. This animal produces B cells that secrete fully human immunoglobulins. The antibodies can be obtained directly from the animal after immunization with an immunogen of interest, as, for example, a preparation of a polyclonal antibody, or alternatively from immortalized B cells derived from the animal, such as hybridomas producing monoclonal antibodies. Additionally, the genes encoding the immunoglobulins with human variable regions can be recovered and expressed to obtain the antibodies directly, or can be further modified to obtain analogs of antibodies such as, for example, single chain Fv molecules.

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An example of a method of producing a nonhuman host, exemplified as a mouse, lacking expression of an endogenous immunoglobulin heavy chain is disclosed in U.S. Patent No. 5,939,598. It can be obtained by a method including deleting the J segment genes from at least one endogenous heavy chain locus in an embryonic stem cell to prevent rearrangement of the locus and to prevent formation of a transcript of a rearranged immunoglobulin heavy chain locus, the deletion being effected by a targeting vector containing a gene encoding a selectable marker; and producing from the embryonic stem cell a transgenic mouse whose somatic and germ cells contain the gene encoding the selectable marker.

A method for producing an antibody of interest, such as a human antibody, is disclosed in U.S. Patent No. 5,916,771. It includes introducing an expression vector that contains a nucleotide sequence encoding a heavy chain into one mammalian host cell in

culture, introducing an expression vector containing a nucleotide sequence encoding a light chain into another mammalian host cell, and fusing the two cells to form a hybrid cell. The hybrid cell expresses an antibody containing the heavy chain and the light chain.

In a further improvement on this procedure, a method for identifying a clinically relevant epitope on an immunogen, and a correlative method for selecting an antibody that binds immunospecifically to the relevant epitope with high affinity, are disclosed in PCT publication WO 99/53049.

4.13.5 FAB FRAGMENTS AND SINGLE CHAIN ANTIBODIES

According to the invention, techniques can be adapted for the production of single-chain antibodies specific to an antigenic protein of the invention (see e.g., U.S. Patent No. 4,946,778). In addition, methods can be adapted for the construction of F_{ab} expression libraries (see e.g., Huse, et al., 1989 Science 246: 1275-1281) to allow rapid and effective identification of monoclonal F_{ab} fragments with the desired specificity for a protein or derivatives, fragments, analogs or homologs thereof. Antibody fragments that contain the idiotypes to a protein antigen may be produced by techniques known in the art including, but not limited to: (i) an $F_{(ab)2}$ fragment produced by pepsin digestion of an antibody molecule; (ii) an F_{ab} fragment generated by reducing the disulfide bridges of an $F_{(ab)2}$ fragment; (iii) an F_{ab} fragment generated by the treatment of the antibody molecule with papain and a reducing agent and (iv) F_{v} fragments.

4.13.6 BISPECIFIC ANTIBODIES

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Bispecific antibodies are monoclonal, preferably human or humanized, antibodies that have binding specificities for at least two different antigens. In the present case, one of the binding specificities is for an antigenic protein of the invention. The second binding target is any other antigen, and advantageously is a cell-surface protein or receptor or receptor subunit.

Methods for making bispecific antibodies are known in the art. Traditionally, the recombinant production of bispecific antibodies is based on the co-expression of two immunoglobulin heavy-chain/light-chain pairs, where the two heavy chains have

different specificities (Milstein and Cuello, Nature, 305:537-539 (1983)). Because of the random assortment of immunoglobulin heavy and light chains, these hybridomas (quadromas) produce a potential mixture of ten different antibody molecules, of which only one has the correct bispecific structure. The purification of the correct molecule is usually accomplished by affinity chromatography steps. Similar procedures are disclosed in WO 93/08829, published 13 May 1993, and in Traunecker et al., 1991 EMBO J., 10:3655-3659.

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Antibody variable domains with the desired binding specificities (antibody-antigen combining sites) can be fused to immunoglobulin constant domain sequences. The fusion preferably is with an immunoglobulin heavy-chain constant domain, comprising at least part of the hinge, CH2, and CH3 regions. It is preferred to have the first heavy-chain constant region (CH1) containing the site necessary for light-chain binding present in at least one of the fusions. DNAs encoding the immunoglobulin heavy-chain fusions and, if desired, the immunoglobulin light chain, are inserted into separate expression vectors, and are co-transfected into a suitable host organism. For further details of generating bispecific antibodies see, for example, Suresh et al., Methods in Enzymology, 121:210 (1986).

According to another approach described in WO 96/27011, the interface between a pair of antibody molecules can be engineered to maximize the percentage of heterodimers that are recovered from recombinant cell culture. The preferred interface comprises at least a part of the CH3 region of an antibody constant domain. In this method, one or more small amino acid side chains from the interface of the first antibody molecule are replaced with larger side chains (e.g. tyrosine or tryptophan). Compensatory "cavities" of identical or similar size to the large side chain(s) are created on the interface of the second antibody molecule by replacing large amino acid side chains with smaller ones (e.g. alanine or threonine). This provides a mechanism for increasing the yield of the heterodimer over other unwanted end-products such as homodimers.

Bispecific antibodies can be prepared as full length antibodies or antibody fragments (e.g. F(ab')₂ bispecific antibodies). Techniques for generating bispecific antibodies from antibody fragments have been described in the literature. For example,

bispecific antibodies can be prepared using chemical linkage. Brennan et al., Science 229:81 (1985) describe a procedure wherein intact antibodies are proteolytically cleaved to generate F(ab')₂ fragments. These fragments are reduced in the presence of the dithiol complexing agent sodium arsenite to stabilize vicinal dithiols and prevent intermolecular disulfide formation. The Fab' fragments generated are then converted to thionitrobenzoate (TNB) derivatives. One of the Fab'-TNB derivatives is then reconverted to the Fab'-thiol by reduction with mercaptoethylamine and is mixed with an equimolar amount of the other Fab'-TNB derivative to form the bispecific antibody. The bispecific antibodies produced can be used as agents for the selective immobilization of enzymes.

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Additionally, Fab' fragments can be directly recovered from E. coli and chemically coupled to form bispecific antibodies. Shalaby et al., <u>J. Exp. Med.</u>
175:217-225 (1992) describe the production of a fully humanized bispecific antibody F(ab')₂ molecule. Each Fab' fragment was separately secreted from E. coli and subjected to directed chemical coupling in vitro to form the bispecific antibody. The bispecific antibody thus formed was able to bind to cells overexpressing the ErbB2 receptor and normal human T cells, as well as trigger the lytic activity of human cytotoxic lymphocytes against human breast tumor targets.

Various techniques for making and isolating bispecific antibody fragments directly from recombinant cell culture have also been described. For example, bispecific antibodies have been produced using leucine zippers. Kostelny et al., <u>J. Immunol.</u>

148(5):1547-1553 (1992). The leucine zipper peptides from the Fos and Jun proteins were linked to the Fab' portions of two different antibodies by gene fusion. The antibody homodimers were reduced at the hinge region to form monomers and then re-oxidized to form the antibody heterodimers. This method can also be utilized for the production of antibody homodimers. The "diabody" technology described by Hollinger et al., <u>Proc. Natl. Acad. Sci. USA</u> 90:6444-6448 (1993) has provided an alternative mechanism for making bispecific antibody fragments. The fragments comprise a heavy-chain variable domain (V_H) connected to a light-chain variable domain (V_L) by a linker which is too short to allow pairing between the two domains on the same chain. Accordingly, the V_H and V_L domains of one fragment are forced to pair with the complementary V_L and V_H

domains of another fragment, thereby forming two antigen-binding sites. Another strategy for making bispecific antibody fragments by the use of single-chain Fv (sFv) dimers has also been reported. See, Gruber et al., J. Immunol. 152:5368 (1994).

Antibodies with more than two valencies are contemplated. For example, trispecific antibodies can be prepared. Tutt et al., J. Immunol. 147:60 (1991).

Exemplary bispecific antibodies can bind to two different epitopes, at least one of which originates in the protein antigen of the invention. Alternatively, an anti-antigenic arm of an immunoglobulin molecule can be combined with an arm which binds to a triggering molecule on a leukocyte such as a T-cell receptor molecule (e.g. CD2, CD3, CD28, or B7), or Fc receptors for IgG (FcyR), such as FcyRI (CD64), FcyRII (CD32) and FcyRII (CD16) so as to focus cellular defense mechanisms to the cell expressing the particular antigen. Bispecific antibodies can also be used to direct cytotoxic agents to cells which express a particular antigen. These antibodies possess an antigen-binding arm and an arm which binds a cytotoxic agent or a radionuclide chelator, such as EOTUBE, DPTA, DOTA, or TETA. Another bispecific antibody of interest binds the protein antigen described herein and further binds tissue factor (TF).

4.13.7 HETEROCONJUGATE ANTIBODIES

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Heteroconjugate antibodies are also within the scope of the present invention. Heteroconjugate antibodies are composed of two covalently joined antibodies. Such antibodies have, for example, been proposed to target immune system cells to unwanted cells (U.S. Patent No. 4,676,980), and for treatment of HIV infection (WO 91/00360; WO 92/200373; EP 03089). It is contemplated that the antibodies can be prepared in vitro using known methods in synthetic protein chemistry, including those involving crosslinking agents. For example, immunotoxins can be constructed using a disulfide exchange reaction or by forming a thioether bond. Examples of suitable reagents for this purpose include iminothiolate and methyl-4-mercaptobutyrimidate and those disclosed, for example, in U.S. Patent No. 4,676,980.

4.13.8 EFFECTOR FUNCTION ENGINEERING

It can be desirable to modify the antibody of the invention with respect to effector function, so as to enhance, e.g., the effectiveness of the antibody in treating cancer. For example, cysteine residue(s) can be introduced into the Fc region, thereby allowing interchain disulfide bond formation in this region. The homodimeric antibody thus generated can have improved internalization capability and/or increased complement-mediated cell killing and antibody-dependent cellular cytotoxicity (ADCC). See Caron et al., J. Exp Med., 176: 1191-1195 (1992) and Shopes, J. Immunol., 148: 2918-2922 (1992). Homodimeric antibodies with enhanced anti-tumor activity can also be prepared using heterobifunctional cross-linkers as described in Wolff et al. Cancer Research, 53: 2560-2565 (1993). Alternatively, an antibody can be engineered that has dual Fc regions and can thereby have enhanced complement lysis and ADCC capabilities. See Stevenson et al., Anti-Cancer Drug Design, 3: 219-230 (1989).

4.13.9 IMMUNOCONJUGATES

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The invention also pertains to immunoconjugates comprising an antibody conjugated to a cytotoxic agent such as a chemotherapeutic agent, toxin (e.g., an enzymatically active toxin of bacterial, fungal, plant, or animal origin, or fragments thereof), or a radioactive isotope (i.e., a radioconjugate).

Chemotherapeutic agents useful in the generation of such immunoconjugates have been described above. Enzymatically active toxins and fragments thereof that can be used include diphtheria A chain, nonbinding active fragments of diphtheria toxin, exotoxin A chain (from Pseudomonas aeruginosa), ricin A chain, abrin A chain, modeccin A chain, alpha-sarcin, Aleurites fordii proteins, dianthin proteins, Phytolaca americana proteins (PAPI, PAPII, and PAP-S), momordica charantia inhibitor, curcin, crotin, sapaonaria officinalis inhibitor, gelonin, mitogellin, restrictocin, phenomycin, enomycin, and the tricothecenes. A variety of radionuclides are available for the production of radioconjugated antibodies. Examples include ²¹²Bi, ¹³¹I, ¹³¹In, ⁹⁰Y, and ¹⁸⁶Re.

Conjugates of the antibody and cytotoxic agent are made using a variety of bifunctional protein-coupling agents such as N-succinimidyl-3-(2-pyridyldithiol) propionate (SPDP), iminothiolane (IT), bifunctional derivatives of imidoesters (such as

dimethyl adipimidate HCL), active esters (such as disuccinimidyl suberate), aldehydes (such as glutareldehyde), bis-azido compounds (such as bis (p-azidobenzoyl) hexanediamine), bis-diazonium derivatives (such as bis-(p-diazoniumbenzoyl)-ethylenediamine), diisocyanates (such as tolyene 2,6-diisocyanate), and bis-active fluorine compounds (such as 1,5-difluoro-2,4-dinitrobenzene). For example, a ricin immunotoxin can be prepared as described in Vitetta et al., Science, 238: 1098 (1987). Carbon-14-labeled 1-isothiocyanatobenzyl-3-methyldiethylene triaminepentaacetic acid (MX-DTPA) is an exemplary chelating agent for conjugation of radionucleotide to the antibody. See WO94/11026.

In another embodiment, the antibody can be conjugated to a "receptor" (such streptavidin) for utilization in tumor pretargeting wherein the antibody-receptor conjugate is administered to the patient, followed by removal of unbound conjugate from the circulation using a clearing agent and then administration of a "ligand" (e.g., avidin) that is in turn conjugated to a cytotoxic agent.

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4.14 COMPUTER READABLE SEQUENCES

In one application of this embodiment, a nucleotide sequence of the present invention can be recorded on computer readable media. As used herein, "computer readable media" refers to any medium which can be read and accessed directly by a computer. Such media include, but are not limited to: magnetic storage media, such as floppy discs, hard disc storage medium, and magnetic tape; optical storage media such as CD-ROM; electrical storage media such as RAM and ROM; and hybrids of these categories such as magnetic/optical storage media. A skilled artisan can readily appreciate how any of the presently known computer readable mediums can be used to create a manufacture comprising computer readable medium having recorded thereon a nucleotide sequence of the present invention. As used herein, "recorded" refers to a process for storing information on computer readable medium. A skilled artisan can readily adopt any of the presently known methods for recording information on computer readable medium to generate manufactures comprising the nucleotide sequence information of the present invention.

A variety of data storage structures are available to a skilled artisan for creating a computer readable medium having recorded thereon a nucleotide sequence of the present invention. The choice of the data storage structure will generally be based on the means chosen to access the stored information. In addition, a variety of data processor programs and formats can be used to store the nucleotide sequence information of the present invention on computer readable medium. The sequence information can be represented in a word processing text file, formatted in commercially-available software such as WordPerfect and Microsoft Word, or represented in the form of an ASCII file, stored in a database application, such as DB2, Sybase, Oracle, or the like. A skilled artisan can readily adapt any number of data processor structuring formats (e.g. text file or database) in order to obtain computer readable medium having recorded thereon the nucleotide sequence information of the present invention.

By providing any of the nucleotide sequences SEQ ID NOs: 1 - 438 or a representative fragment thereof; or a nucleotide sequence at least 95% identical to any of the nucleotide sequences of SEQ ID NOs: 1 - 438 in computer readable form, a skilled artisan can routinely access the sequence information for a variety of purposes.

Computer software is publicly available which allows a skilled artisan to access sequence information provided in a computer readable medium. The examples which follow demonstrate how software which implements the BLAST (Altschul et al., J. Mol. Biol. 215:403-410 (1990)) and BLAZE (Brutlag et al., Comp. Chem. 17:203-207 (1993)) search algorithms on a Sybase system is used to identify open reading frames (ORFs) within a nucleic acid sequence. Such ORFs may be protein encoding fragments and may be useful in producing commercially important proteins such as enzymes used in fermentation reactions and in the production of commercially useful metabolites.

As used herein, "a computer-based system" refers to the hardware means, software means, and data storage means used to analyze the nucleotide sequence information of the present invention. The minimum hardware means of the computer-based systems of the present invention comprises a central processing unit (CPU), input means, output means, and data storage means. A skilled artisan can readily appreciate that any one of the currently available computer-based systems are suitable for use in the present invention. As stated above, the computer-based systems of the present

invention comprise a data storage means having stored therein a nucleotide sequence of the present invention and the necessary hardware means and software means for supporting and implementing a search means. As used herein, "data storage means" refers to memory which can store nucleotide sequence information of the present invention, or a memory access means which can access manufactures having recorded thereon the nucleotide sequence information of the present invention.

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As used herein, "search means" refers to one or more programs which are implemented on the computer-based system to compare a target sequence or target structural motif with the sequence information stored within the data storage means. Search means are used to identify fragments or regions of a known sequence which match a particular target sequence or target motif. A variety of known algorithms are disclosed publicly and a variety of commercially available software for conducting search means are and can be used in the computer-based systems of the present invention. Examples of such software includes, but is not limited to, Smith-Waterman, MacPattern (EMBL), BLASTN and BLASTA (NPOLYPEPTIDEIA). A skilled artisan can readily recognize that any one of the available algorithms or implementing software packages for conducting homology searches can be adapted for use in the present computer-based systems. As used herein, a "target sequence" can be any nucleic acid or amino acid sequence of six or more nucleotides or two or more amino acids. A skilled artisan can readily recognize that the longer a target sequence is, the less likely a target sequence will be present as a random occurrence in the database. The most preferred sequence length of a target sequence is from about 10 to 300 amino acids, more preferably from about 30 to 100 nucleotide residues. However, it is well recognized that searches for commercially important fragments, such as sequence fragments involved in gene expression and protein processing, may be of shorter length.

As used herein, "a target structural motif," or "target motif," refers to any rationally selected sequence or combination of sequences in which the sequence(s) are chosen based on a three-dimensional configuration which is formed upon the folding of the target motif. There are a variety of target motifs known in the art. Protein target motifs include, but are not limited to, enzyme active sites and signal sequences. Nucleic

acid target motifs include, but are not limited to, promoter sequences, hairpin structures and inducible expression elements (protein binding sequences).

4.15 TRIPLE HELIX FORMATION

5 In addition, the fragments of the present invention, as broadly described, can be used to control gene expression through triple helix formation or antisense DNA or RNA, both of which methods are based on the binding of a polynucleotide sequence to DNA or RNA. Polynucleotides suitable for use in these methods are preferably 20 to 40 bases in length and are designed to be complementary to a region of the gene involved in 10 transcription (triple helix - see Lee et al., Nucl. Acids Res. 6:3073 (1979); Cooney et al., Science 15241:456 (1988); and Dervan et al., Science 251:1360 (1991)) or to the mRNA itself (antisense - Olmno, J. Neurochem. 56:560 (1991); Oligodeoxynucleotides as Antisense Inhibitors of Gene Expression, CRC Press, Boca Raton, FL (1988)). Triple helix-formation optimally results in a shut-off of RNA transcription from DNA, while 15 antisense RNA hybridization blocks translation of an mRNA molecule into polypeptide. Both techniques have been demonstrated to be effective in model systems. Information contained in the sequences of the present invention is necessary for the design of an antisense or triple helix oligonucleotide.

4.16 DIAGNOSTIC ASSAYS AND KITS

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The present invention further provides methods to identify the presence or expression of one of the ORFs of the present invention, or homolog thereof, in a test sample, using a nucleic acid probe or antibodies of the present invention, optionally conjugated or otherwise associated with a suitable label.

In general, methods for detecting a polynucleotide of the invention can comprise contacting a sample with a compound that binds to and forms a complex with the polynucleotide for a period sufficient to form the complex, and detecting the complex, so that if a complex is detected, a polynucleotide of the invention is detected in the sample. Such methods can also comprise contacting a sample under stringent hybridization conditions with nucleic acid primers that anneal to a polynucleotide of the invention

under such conditions, and amplifying annealed polynucleotides, so that if a polynucleotide is amplified, a polynucleotide of the invention is detected in the sample.

In general, methods for detecting a polypeptide of the invention can comprise contacting a sample with a compound that binds to and forms a complex with the polypeptide for a period sufficient to form the complex, and detecting the complex, so that if a complex is detected, a polypeptide of the invention is detected in the sample.

In detail, such methods comprise incubating a test sample with one or more of the antibodies or one or more of the nucleic acid probes of the present invention and assaying for binding of the nucleic acid probes or antibodies to components within the test sample.

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Conditions for incubating a nucleic acid probe or antibody with a test sample vary. Incubation conditions depend on the format employed in the assay, the detection methods employed, and the type and nature of the nucleic acid probe or antibody used in the assay. One skilled in the art will recognize that any one of the commonly available hybridization, amplification or immunological assay formats can readily be adapted to employ the nucleic acid probes or antibodies of the present invention. Examples of such assays can be found in Chard, T., An Introduction to Radioimmunoassay and Related Techniques, Elsevier Science Publishers, Amsterdam, The Netherlands (1986); Bullock, G.R. et al., Techniques in Immunocytochemistry, Academic Press, Orlando, FL Vol. 1 (1982), Vol. 2 (1983), Vol. 3 (1985); Tijssen, P., Practice and Theory of immunoassays: Laboratory Techniques in Biochemistry and Molecular Biology, Elsevier Science Publishers, Amsterdam, The Netherlands (1985). The test samples of the present invention include cells, protein or membrane extracts of cells, or biological fluids such as sputum, blood, serum, plasma, or urine. The test sample used in the above-described method will vary based on the assay format, nature of the detection method and the tissues, cells or extracts used as the sample to be assayed. Methods for preparing protein extracts or membrane extracts of cells are well known in the art and can be readily be adapted in order to obtain a sample which is compatible with the system utilized.

In another embodiment of the present invention, kits are provided which contain the necessary reagents to carry out the assays of the present invention. Specifically, the invention provides a compartment kit to receive, in close confinement, one or more containers which comprises: (a) a first container comprising one of the probes or

antibodies of the present invention; and (b) one or more other containers comprising one or more of the following: wash reagents, reagents capable of detecting presence of a bound probe or antibody.

In detail, a compartment kit includes any kit in which reagents are contained in separate containers. Such containers include small glass containers, plastic containers or strips of plastic or paper. Such containers allows one to efficiently transfer reagents from one compartment to another compartment such that the samples and reagents are not cross-contaminated, and the agents or solutions of each container can be added in a quantitative fashion from one compartment to another. Such containers will include a container which will accept the test sample, a container which contains the antibodies used in the assay, containers which contain wash reagents (such as phosphate buffered saline, Tris-buffers, etc.), and containers which contain the reagents used to detect the bound antibody or probe. Types of detection reagents include labeled nucleic acid probes, labeled secondary antibodies, or in the alternative, if the primary antibody is labeled, the enzymatic, or antibody binding reagents which are capable of reacting with the labeled antibody. One skilled in the art will readily recognize that the disclosed probes and antibodies of the present invention can be readily incorporated into one of the established kit formats which are well known in the art.

4.17 MEDICAL IMAGING

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The novel polypeptides and binding partners of the invention are useful in medical imaging of sites expressing the molecules of the invention (e.g., where the polypeptide of the invention is involved in the immune response, for imaging sites of inflammation or infection). See, e.g., Kunkel et al., U.S. Pat. NO. 5,413,778. Such methods involve chemical attachment of a labeling or imaging agent, administration of the labeled polypeptide to a subject in a pharmaceutically acceptable carrier, and imaging the labeled polypeptide *in vivo* at the target site.

4.18 SCREENING ASSAYS

Using the isolated proteins and polynucleotides of the invention, the present invention further provides methods of obtaining and identifying agents which bind to a

polypeptide encoded by an ORF corresponding to any of the nucleotide sequences set forth in SEQ ID NOs: 1 - 438, or bind to a specific domain of the polypeptide encoded by the nucleic acid. In detail, said method comprises the steps of:

(a) contacting an agent with an isolated protein encoded by an ORF of the present invention, or nucleic acid of the invention; and

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(b) determining whether the agent binds to said protein or said nucleic acid.

In general, therefore, such methods for identifying compounds that bind to a polynucleotide of the invention can comprise contacting a compound with a polynucleotide of the invention for a time sufficient to form a polynucleotide/compound complex, and detecting the complex, so that if a polynucleotide/compound complex is detected, a compound that binds to a polynucleotide of the invention is identified.

Likewise, in general, therefore, such methods for identifying compounds that bind to a polypeptide of the invention can comprise contacting a compound with a polypeptide of the invention for a time sufficient to form a polypeptide/compound complex, and detecting the complex, so that if a polypeptide/compound complex is detected, a compound that binds to a polynucleotide of the invention is identified.

Methods for identifying compounds that bind to a polypeptide of the invention can also comprise contacting a compound with a polypeptide of the invention in a cell for a time sufficient to form a polypeptide/compound complex, wherein the complex drives expression of a receptor gene sequence in the cell, and detecting the complex by detecting reporter gene sequence expression, so that if a polypeptide/compound complex is detected, a compound that binds a polypeptide of the invention is identified.

Compounds identified via such methods can include compounds which modulate the activity of a polypeptide of the invention (that is, increase or decrease its activity, relative to activity observed in the absence of the compound). Alternatively, compounds identified via such methods can include compounds which modulate the expression of a polynucleotide of the invention (that is, increase or decrease expression relative to expression levels observed in the absence of the compound). Compounds, such as compounds identified via the methods of the invention, can be tested using standard assays well known to those of skill in the art for their ability to modulate activity/expression.

The agents screened in the above assay can be, but are not limited to, peptides, carbohydrates, vitamin derivatives, or other pharmaceutical agents. The agents can be selected and screened at random or rationally selected or designed using protein modeling techniques.

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For random screening, agents such as peptides, carbohydrates, pharmaceutical agents and the like are selected at random and are assayed for their ability to bind to the protein encoded by the ORF of the present invention. Alternatively, agents may be rationally selected or designed. As used herein, an agent is said to be "rationally selected or designed" when the agent is chosen based on the configuration of the particular protein. For example, one skilled in the art can readily adapt currently available procedures to generate peptides, pharmaceutical agents and the like, capable of binding to a specific peptide sequence, in order to generate rationally designed antipeptide peptides, for example see Hurby et al., Application of Synthetic Peptides: Antisense Peptides," In Synthetic Peptides, A User's Guide, W.H. Freeman, NY (1992), pp. 289-307, and Kaspczak et al., Biochemistry 28:9230-8 (1989), or pharmaceutical agents, or the like.

In addition to the foregoing, one class of agents of the present invention, as broadly described, can be used to control gene expression through binding to one of the ORFs or EMFs of the present invention. As described above, such agents can be randomly screened or rationally designed/selected. Targeting the ORF or EMF allows a skilled artisan to design sequence specific or element specific agents, modulating the expression of either a single ORF or multiple ORFs which rely on the same EMF for expression control. One class of DNA binding agents are agents which contain base residues which hybridize or form a triple helix formation by binding to DNA or RNA. Such agents can be based on the classic phosphodiester, ribonucleic acid backbone, or can be a variety of sulfhydryl or polymeric derivatives which have base attachment capacity.

Agents suitable for use in these methods preferably contain 20 to 40 bases and are designed to be complementary to a region of the gene involved in transcription (triple helix - see Lee et al., Nucl. Acids Res. 6:3073 (1979); Cooney et al., Science 241:456 (1988); and Dervan et al., Science 251:1360 (1991)) or to the mRNA itself (antisense - Okano, J. Neurochem. 56:560 (1991); Oligodeoxynucleotides as Antisense Inhibitors of

Gene Expression, CRC Press, Boca Raton, FL (1988)). Triple helix-formation optimally results in a shut-off of RNA transcription from DNA, while antisense RNA hybridization blocks translation of an mRNA molecule into polypeptide. Both techniques have been demonstrated to be effective in model systems. Information contained in the sequences of the present invention is necessary for the design of an antisense or triple helix oligonucleotide and other DNA binding agents.

Agents which bind to a protein encoded by one of the ORFs of the present invention can be used as a diagnostic agent. Agents which bind to a protein encoded by one of the ORFs of the present invention can be formulated using known techniques to generate a pharmaceutical composition.

4.19 USE OF NUCLEIC ACIDS AS PROBES

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Another aspect of the subject invention is to provide for polypeptide-specific nucleic acid hybridization probes capable of hybridizing with naturally occurring nucleotide sequences. The hybridization probes of the subject invention may be derived from any of the nucleotide sequences SEQ ID NOs: 1 - 438. Because the corresponding gene is only expressed in a limited number of tissues, a hybridization probe derived from of any of the nucleotide sequences SEQ ID NOs: 1 - 438 can be used as an indicator of the presence of RNA of cell type of such a tissue in a sample.

Any suitable hybridization technique can be employed, such as, for example, in situ hybridization. PCR as described in US Patents Nos. 4,683,195 and 4,965,188 provides additional uses for oligonucleotides based upon the nucleotide sequences. Such probes used in PCR may be of recombinant origin, may be chemically synthesized, or a mixture of both. The probe will comprise a discrete nucleotide sequence for the detection of identical sequences or a degenerate pool of possible sequences for identification of closely related genomic sequences.

Other means for producing specific hybridization probes for nucleic acids include the cloning of nucleic acid sequences into vectors for the production of mRNA probes. Such vectors are known in the art and are commercially available and may be used to synthesize RNA probes *in vitro* by means of the addition of the appropriate RNA polymerase as T7 or SP6 RNA polymerase and the appropriate radioactively labeled

nucleotides. The nucleotide sequences may be used to construct hybridization probes for mapping their respective genomic sequences. The nucleotide sequence provided herein may be mapped to a chromosome or specific regions of a chromosome using well known genetic and/or chromosomal mapping techniques. These techniques include in situ hybridization, linkage analysis against known chromosomal markers, hybridization screening with libraries or flow-sorted chromosomal preparations specific to known chromosomes, and the like. The technique of fluorescent in situ hybridization of chromosome spreads has been described, among other places, in Verma et al (1988) Human Chromosomes: A Manual of Basic Techniques, Pergamon Press, New York NY.

Fluorescent in situ hybridization of chromosomal preparations and other physical chromosome mapping techniques may be correlated with additional genetic map data. Examples of genetic map data can be found in the 1994 Genome Issue of Science (265:1981f). Correlation between the location of a nucleic acid on a physical chromosomal map and a specific disease (or predisposition to a specific disease) may help delimit the region of DNA associated with that genetic disease. The nucleotide sequences of the subject invention may be used to detect differences in gene sequences between normal, carrier or affected individuals.

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4.20 PREPARATION OF SUPPORT BOUND OLIGONUCLEOTIDES

Support bound oligonucleotides may be prepared by any of the methods known to

Oligonucleotides, i.e., small nucleic acid segments, may be readily prepared by, for example, directly synthesizing the oligonucleotide by chemical means, as is commonly practiced using an automated oligonucleotide synthesizer.

those of skill in the art using any suitable support such as glass, polystyrene or Teflon. One strategy is to precisely spot oligonucleotides synthesized by standard synthesizers. Immobilization can be achieved using passive adsorption (Inouye & Hondo, (1990) J. Clin. Microbiol. 28(6) 1469-72); using UV light (Nagata *et al.*, 1985; Dahlen *et al.*, 1987; Morrissey & Collins, (1989) Mol. Cell Probes 3(2) 189-207) or by covalent binding of base modified DNA (Keller *et al.*, 1988; 1989); all references being specifically incorporated herein.

Another strategy that may be employed is the use of the strong biotin-streptavidin interaction as a linker. For example, Broude et al. (1994) Proc. Natl. Acad. Sci. USA 91(8)

3072-6, describe the use of biotinylated probes, although these are duplex probes, that are immobilized on streptavidin-coated magnetic beads. Streptavidin-coated beads may be purchased from Dynal, Oslo. Of course, this same linking chemistry is applicable to coating any surface with streptavidin. Biotinylated probes may be purchased from various sources, such as, e.g., Operon Technologies (Alameda, CA).

Nunc Laboratories (Naperville, IL) is also selling suitable material that could be used. Nunc Laboratories have developed a method by which DNA can be covalently bound to the microwell surface termed Covalink NH. CovaLink NH is a polystyrene surface grafted with secondary amino groups (>NH) that serve as bridge-heads for further covalent coupling. CovaLink Modules may be purchased from Nunc Laboratories. DNA molecules may be bound to CovaLink exclusively at the 5'-end by a phosphoramidate bond, allowing immobilization of more than 1 pmol of DNA (Rasmussen et al., (1991) Anal. Biochem. 198(1) 138-42).

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The use of CovaLink NH strips for covalent binding of DNA molecules at the 5'-end has been described (Rasmussen et al., (1991). In this technology, a phosphoramidate bond is employed (Chu et al., (1983) Nucleic Acids Res. 11(8) 6513-29). This is beneficial as immobilization using only a single covalent bond is preferred. The phosphoramidate bond joins the DNA to the CovaLink NH secondary amino groups that are positioned at the end of spacer arms covalently grafted onto the polystyrene surface through a 2 nm long spacer arm. To link an oligonucleotide to CovaLink NH via an phosphoramidate bond, the oligonucleotide terminus must have a 5'-end phosphate group. It is, perhaps, even possible for biotin to be covalently bound to CovaLink and then streptavidin used to bind the probes.

More specifically, the linkage method includes dissolving DNA in water (7.5 ng/ul) and denaturing for 10 min. at 95°C and cooling on ice for 10 min. Ice-cold 0.1 M 1-methylimidazole, pH 7.0 (1-MeIm₇), is then added to a final concentration of 10 mM 1-MeIm₇. A ss DNA solution is then dispensed into CovaLink NH strips (75 ul/well) standing on ice.

Carbodiimide 0.2 M 1-ethyl-3-(3-dimethylaminopropyl)-carbodiimide (EDC), dissolved in 10 mM 1-MeIm₇, is made fresh and 25 ul added per well. The strips are incubated for 5 hours at 50°C. After incubation the strips are washed using, e.g., Nunc-Immuno Wash; first the wells are washed 3 times, then they are soaked with washing

solution for 5 min., and finally they are washed 3 times (where in the washing solution is 0.4 N NaOH, 0.25% SDS heated to 50°C).

It is contemplated that a further suitable method for use with the present invention is that described in PCT Patent Application WO 90/03382 (Southern & Maskos), incorporated herein by reference. This method of preparing an oligonucleotide bound to a support involves attaching a nucleoside 3'-reagent through the phosphate group by a covalent phosphodiester link to aliphatic hydroxyl groups carried by the support. The oligonucleotide is then synthesized on the supported nucleoside and protecting groups removed from the synthetic oligonucleotide chain under standard conditions that do not cleave the oligonucleotide from the support. Suitable reagents include nucleoside phosphoramidite and nucleoside hydrogen phosphorate.

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An on-chip strategy for the preparation of DNA probe for the preparation of DNA probe arrays may be employed. For example, addressable laser-activated photodeprotection may be employed in the chemical synthesis of oligonucleotides directly on a glass surface, as described by Fodor et al. (1991) Science 251(4995) 767-73, incorporated herein by reference. Probes may also be immobilized on nylon supports as described by Van Ness et al. (1991) Nucleic Acids Res. 19(12) 3345-50; or linked to Teflon using the method of Duncan & Cavalier (1988) Anal. Biochem. 169(1) 104-8; all references being specifically incorporated herein.

To link an oligonucleotide to a nylon support, as described by Van Ness et al. (1991), requires activation of the nylon surface via alkylation and selective activation of the 5'-amine of oligonucleotides with cyanuric chloride.

One particular way to prepare support bound oligonucleotides is to utilize the light-generated synthesis described by Pease et al., (1994) PNAS USA 91(11) 5022-6, incorporated herein by reference). These authors used current photolithographic techniques to generate arrays of immobilized oligonucleotide probes (DNA chips). These methods, in which light is used to direct the synthesis of oligonucleotide probes in high-density, miniaturized arrays, utilize photolabile 5'-protected N-acyl-deoxynucleoside phosphoramidites, surface linker chemistry and versatile combinatorial synthesis strategies.

30 A matrix of 256 spatially defined oligonucleotide probes may be generated in this manner.

4.21 PREPARATION OF NUCLEIC ACID FRAGMENTS

The nucleic acids may be obtained from any appropriate source, such as cDNAs, genomic DNA, chromosomal DNA, microdissected chromosome bands, cosmid or YAC inserts, and RNA, including mRNA without any amplification steps. For example, Sambrook *et al.* (1989) describes three protocols for the isolation of high molecular weight DNA from mammalian cells (p. 9.14-9.23).

DNA fragments may be prepared as clones in M13, plasmid or lambda vectors and/or prepared directly from genomic DNA or cDNA by PCR or other amplification methods. Samples may be prepared or dispensed in multiwell plates. About 100-1000 ng of DNA samples may be prepared in 2-500 ml of final volume.

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The nucleic acids would then be fragmented by any of the methods known to those of skill in the art including, for example, using restriction enzymes as described at 9.24-9.28 of Sambrook *et al.* (1989), shearing by ultrasound and NaOH treatment.

Low pressure shearing is also appropriate, as described by Schriefer *et al.* (1990) Nucleic Acids Res. 18(24) 7455-6, incorporated herein by reference). In this method, DNA samples are passed through a small French pressure cell at a variety of low to intermediate pressures. A lever device allows controlled application of low to intermediate pressures to the cell. The results of these studies indicate that low-pressure shearing is a useful alternative to sonic and enzymatic DNA fragmentation methods.

One particularly suitable way for fragmenting DNA is contemplated to be that using the two base recognition endonuclease, CviJI, described by Fitzgerald et al. (1992) Nucleic Acids Res. 20(14) 3753-62. These authors described an approach for the rapid fragmentation and fractionation of DNA into particular sizes that they contemplated to be suitable for shotgun cloning and sequencing.

The restriction endonuclease CviJI normally cleaves the recognition sequence PuGCPy between the G and C to leave blunt ends. Atypical reaction conditions, which alter the specificity of this enzyme (CviJI**), yield a quasi-random distribution of DNA fragments form the small molecule pUC19 (2688 base pairs). Fitzgerald et al. (1992) quantitatively evaluated the randomness of this fragmentation strategy, using a CviJI** digest of pUC19 that was size fractionated by a rapid gel filtration method and directly ligated, without end repair, to a lac Z minus M13 cloning vector. Sequence analysis of 76

clones showed that $Cvi\Pi^{**}$ restricts pyGCPy and PuGCPu, in addition to PuGCPy sites, and that new sequence data is accumulated at a rate consistent with random fragmentation.

As reported in the literature, advantages of this approach compared to sonication and agarose gel fractionation include: smaller amounts of DNA are required (0.2-0.5 ug instead of 2-5 ug); and fewer steps are involved (no preligation, end repair, chemical extraction, or agarose gel electrophoresis and elution are needed

Irrespective of the manner in which the nucleic acid fragments are obtained or prepared, it is important to denature the DNA to give single stranded pieces available for hybridization. This is achieved by incubating the DNA solution for 2-5 minutes at 80-90°C. The solution is then cooled quickly to 2°C to prevent renaturation of the DNA fragments before they are contacted with the chip. Phosphate groups must also be removed from genomic DNA by methods known in the art.

4.22 PREPARATION OF DNA ARRAYS

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Arrays may be prepared by spotting DNA samples on a support such as a nylon membrane. Spotting may be performed by using arrays of metal pins (the positions of which correspond to an array of wells in a microtiter plate) to repeated by transfer of about 20 nl of a DNA solution to a nylon membrane. By offset printing, a density of dots higher than the density of the wells is achieved. One to 25 dots may be accommodated in 1 mm², depending on the type of label used. By avoiding spotting in some preselected number of rows and columns, separate subsets (subarrays) may be formed. Samples in one subarray may be the same genomic segment of DNA (or the same gene) from different individuals, or may be different, overlapped genomic clones. Each of the subarrays may represent replica spotting of the same samples. In one example, a selected gene segment may be amplified from 64 patients. For each patient, the amplified gene segment may be in one 96-well plate (all 96 wells containing the same sample). A plate for each of the 64 patients is prepared. By using a 96-pin device, all samples may be spotted on one 8 x 12 cm membrane. Subarrays may contain 64 samples, one from each patient. Where the 96 subarrays are identical, the dot span may be 1 mm² and there may be a 1 mm space between subarrays.

Another approach is to use membranes or plates (available from NUNC, Naperville, Illinois) which may be partitioned by physical spacers e.g. a plastic grid molded over the membrane, the grid being similar to the sort of membrane applied to the bottom of multiwell

plates, or hydrophobic strips. A fixed physical spacer is not preferred for imaging by exposure to flat phosphor-storage screens or x-ray films.

The present invention is illustrated in the following examples. Upon consideration of the present disclosure, one of skill in the art will appreciate that many other embodiments and variations may be made in the scope of the present invention. Accordingly, it is intended that the broader aspects of the present invention not be limited to the disclosure of the following examples. The present invention is not to be limited in scope by the exemplified embodiments which are intended as illustrations of single aspects of the invention, and compositions and methods which are functionally equivalent are within the scope of the invention. Indeed, numerous modifications and variations in the practice of the invention are expected to occur to those skilled in the art upon consideration of the present preferred embodiments. Consequently, the only limitations which should be placed upon the scope of the invention are those which appear in the appended claims.

All references cited within the body of the instant specification are hereby incorporated by reference in their entirety.

5.0 EXAMPLES

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5.1 EXAMPLE 1

Novel Nucleic Acid Sequences Obtained From Various Libraries

A plurality of novel nucleic acids were obtained from cDNA libraries prepared from various human tissues and in some cases isolated from a genomic library derived from human chromosome using standard PCR, SBH sequence signature analysis and Sanger sequencing techniques. The inserts of the library were amplified with PCR using primers specific for the vector sequences which flank the inserts. Clones from cDNA libraries were spotted on nylon membrane filters and screened with oligonucleotide probes (e.g., 7-mers) to obtain signature sequences. The clones were clustered into groups of similar or identical sequences. Representative clones were selected for sequencing.

In some cases, the 5' sequence of the amplified inserts was then deduced using a typical Sanger sequencing protocol. PCR products were purified and subjected to fluorescent dye terminator cycle sequencing. Single pass gel sequencing was done using a 377 Applied Biosystems (ABI) sequencer to obtain the novel nucleic acid sequences. In

some cases RACE (Random Amplification of cDNA Ends) was performed to further extend the sequence in the 5' direction.

5.2 EXAMPLE 2

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Novel Nucleic Acids

The novel nucleic acids of the present invention of the invention were assembled from sequences that were obtained from a cDNA library by methods described in Example 1 above, and in some cases sequences obtained from one or more public databases. The nucleic acids were assembled using an EST sequence as a seed. Then a recursive algorithm was used to extend the seed EST into an extended assemblage, by pulling additional sequences from different databases (i.e., Hyseq's database containing EST sequences, dbEST version 119, gb pri 119, and UniGene version 119) that belong to this assemblage. The algorithm terminated when there was no additional sequences from the above databases that would extend the assemblage. Inclusion of component sequences into the assemblage was based on a BLASTN hit to the extending assemblage with BLAST score greater than 300 and percent identity greater than 95%.

Using PHRAP (Univ. of Washington) or CAP4 (Paracel), a full length gene cDNA sequence and its corresponding protein sequence were generated from the assemblage. Any frame shifts and incorrect stop codons were corrected by hand editing. During editing, the sequence was checked using FASTY and/or BLAST against Genbank (i.e., dbEST version 120, gb pri 120, UniGene version 120, Genpept release 120). Other computer programs which may have been used in the editing process were phredPhrap and Consed (University of Washington) and ed-ready, ed-ext and cg-zip-2 (Hyseq, Inc.). The full-length nucleotide and amino acid sequences, including splice variants resulting from these procedures are shown in the Sequence Listing as SEQ ID NOS: 1-438.

Table 1 shows the various tissue sources of SEQ ID NO: 1-438.

The nearest neighbor results for polypeptides encoded by SEQ ID NO: 1-438 were obtained by a BLASTP (version 2.0al 19MP-WashU) search against Genpept, Geneseq and SwissProt databases using BLAST algorithm. The nearest neighbor result showed the closest homologue with functional annotation for SEQ ID NO: 1-438. The translated amino acid sequences for which the nucleic acid sequence encodes are shown in the Sequence Listing. The homologues with identifiable functions for SEQ ID NO: 1-

438 are shown in Table 2 below. Using eMatrix software package (Stanford University, Stanford, CA) (Wu et al., J. Comp. Biol., Vol. 6 pp. 219-235 (1999) herein incorporated by reference), all the sequences were examined to determine whether they had identifiable signature regions. Table 3 shows the signature region found in the indicated polypeptide sequences, the description of the signature, the eMatrix p-value(s) and the position(s) of the signature within the polypeptide sequence.

Using the Pfam software program (Sonnhammer et al., Nucleic Acids Res., Vol. 26(1) pp. 320-322 (1998) herein incorporated by reference) polypeptides encoded by SEQ ID NO: 1-438 (i.e. SEQ ID NO: 1-438) were examined for domains with homology to certain peptide domains. Table 4 shows the name of the domain found, the description, the product of all the e-value of similar domains found, the pFam score for the identified domain within the sequence, number of similar domains found, and the position of the domain in the SEQ ID NO: being interrorgated.

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The GeneAtlas™ software package (Molecular Simulations Inc. (MSI), San 15 Diego, CA) was used to predict the three-dimensional structure models for the polypeptides encoded by SEQ ID NO: 1-438 (i.e. SEQ ID NO: 1-438). Models were generated by (1) PSI-BLAST which is a multiple alignment sequence profile-based searching developed by Altschul et al, (Nucl. Acids. Res. 25, 3389-3408 (1997)), (2) High Throughput Modeling (HTM) (Molecular Simulations Inc. (MSI) San Diego, CA,) 20 which is an automated sequence and structure searching procedure (http://www.msi.com/), and (3) SeqFold™ which is a fold recognition method described by Fischer and Eisenberg (J. Mol. Biol. 209, 779-791 (1998)). This analysis was carried out, in part, by comparing the polypeptides of the invention with the known NMR (nuclear magnetic resonance) and x-ray crystal three-dimensional structures as templates. Table 5 shows, "PDB ID", the Protein DataBase (PDB) identifier given to template 25 structure; "Chain ID", identifier of the subcomponent of the PDB template structure; "Compound Information", information of the PDB template structure and/or its subcomponents; "PDB Function Annotation" gives function of the PDB template as annotated by the PDB files (http://www.rcsb.org/PDB/); start and end amino acid position of the protein sequence aligned; PSI-BLAST score, the verify score, the SeqFold score, 30 and the Potential(s) of Mean Force (PMF). The verify score is produced by GeneAtlas™

software (MSI), is based on Dr. Eisenberg's Profile-3D threading program developed in Dr. David Eisenberg's laboratory (US patent no. 5,436,850 and Luthy, Bowie, and Eisenberg, Nature, 356:83-85 (1992)) and a publication by R. Sanchez and A. Sali, Proc. Natl. Acad. Sci. USA, 95:13597-12502. The verify score produced by GeneAtlas normalizes the verify score for proteins with different lengths so that a unified cutoff can be used to select good models as follows:

Verify score (normalized) = (raw score - 1/2 high score)/(1/2 high score)

5

10

15

20

30

The PFM score, produced by GeneAtlas™ software (MSI), is a composite scoring function that depends in part on the compactness of the model, sequence identity in the alignment used to build the model, pairwise and surface mean force potentials (MFP). As given in Table 5, a verify score between 0 to 1.0, with 1 being the best, represents a good model. Similarly, a PMF score between 0 to 1.0, with 1 being the best, represents a good model. A SeqFold™ score of more than 50 is considered significant. A good model may also be determined by one of skill in the art based all the information in Table 5 taken in totality.

The nucleotide sequence within the sequences that codes for signal peptide sequences and their cleavage sites can be determined from using Neural Network SignalP V1.1 program (from Center for Biological Sequence Analysis, The Technical University of Denmark). The process for identifying prokaryotic and eukaryotic signal peptides and their cleavage sites are also disclosed by Henrik Nielson, Jacob Engelbrecht, Soren Brunak, and Gunnar von Heijne in the publication "Identification of prokaryotic and eukaryotic signal peptides and prediction of their cleavage sites" Protein Engineering, Vol. 10, no. 1, pp. 1-6 (1997), incorporated herein by reference. A maximum S score and a mean S score, as described in the Nielson et as reference, was obtained for the polypeptide sequences. Table 6 shows the position of the signal peptide in each of the polypeptides and the maximum score and mean score associated with that signal peptide. Table 7 correlates each of SEQ ID NO: 1-438 to a specific chromosomal location.

Table 8 is a correlation table of the novel polynucleotide sequences SEQ ID NO: 1-438, novel polypeptide sequences SEQ ID NO: 1-438, and their corresponding priority

nucleotide sequences in the priority application USSN 09/774,528, herein incorporated by reference in its entirety.

Table 1

Tissue Origin	RNA/Tissue	Library	SEQ ID NO:
	Source	Name	
adult brain	GIBCO	AB3001	76-77 91 106-107 115 134 163-164 178 203 232 255 265 276 279 322-323
adult brain	GIBCO	ABD003	16 19 24 77 80-81 85 89-90 92 96 98 105
İ	,		110 116 121-123 125 130-132 134-136 138
			142-143 151 153 158-159 163-164 184 191
			193 196 198 200 208-209 213-214 216 219-
			220 223 229 232-234 236 239 241 243 257-
			259 262 265 267 274-276 278 284 292 302
			317 321 324-325 327 337-338 340 348 359 371 391-392 400
adult brain	Clontech	ABR001	1 18-19 35 80 98 125 136 153 185 200 209
			221 228-229 239 243 274-275 302 399-400
adult brain	Clontech	ABR0065	7-8 18 32 35 52 57 85 91 96 111 113 126
•			131 135 138-139 142 148 153-154 181 188
			192 199 209-211 217 221 224 226 229 233 235 238 243 248 273 283-284 286 292 316
	Ì		322 348 357 361 367 376 378 399 407 409
			417 428
adult brain	Clontech	ABR008	2 4 6-11 19-21 23-25 31 35-37 39-41 45-46
1			72-73 76 80-81 85 88-90 94-95 97 102-105
	·		109 111-112 114-119 121-122 126-131 134-
			135 138-139 144 146-150 152-153 156-157
		÷	159 168-172 174-175 178 180 182 185-186
			219 221-222 224 229-230 232-233 236-239
			243-244 248 253-256 260-261 263-265 273
			276 281-282 286-289 291-292 299-300 302
			304 315-317 319 321-322 324 326 329 331-
	}	-	332 341 352-357 360 362 365 367-368 370
			376-377 379-380 383-384 387-389 391-392
	•		394 396-402 407-410 412-413 419 425-426
adult brain	Clontech	ABR011	85 90
adult brain	BioChain	ABR012	148 213
adult brain	BioChain	ABR013	85 322
adult brain	Invitrogen	ABR014	9 23 85 146 200 233 282 321 330
adult brain	Invitrogen	ABR015	14 31 69 121 124 163 209 216 224 291 377
adult brain	Invitrogen	ABR016	92 136 219 279
adult brain	Invitrogen	ABT004	2 7-8 20-21 33 85 90-91 95 97 102-103 108 121 123 129-131 138-139 143 146 151 153
	1		157-158 172 178 180 209-210 213 219 229-
			230 232 234 239 308 321 330 360 365 370-
			373 375 401 412
adipocytes	Stratagene	ADP001	3-4 23 36 79 81 106-107 116 129 133-134
	1		147 151 154 158 179 181 192 196 222 230
admonal miss i	03 cm h = -3:	222000	256-257 287 292 297 313 329 359
adrenal gland	Clontech	ADR002	2 25 27 33 57 76 85-86 88 96 98 105-108 114 121-122 125 129-130 134 147 164 178
			180 182 198-199 201 205 207-208 240-241
			244 246 253-254 257 261 276 280 292 320
			329 336 352 403
adult heart	GIBCO	AHR001	3 17-21 27 32 74 76 85 89-91 95-96 102-103
•			105-110 117 121 124-125 128 131 134-136
	}		139 141 148 151-153 155-156 161 163 181-
			182 186 190 193 198 200-201 205 207 211-
L	L		213 215 222

Table 1

Tissue Origin	RNA/Tissue	Library	SEQ ID NO:
	Source	Name	
			225 229-230 234 251-254 257-259 263 274-
			277 280 292-297 301 303-304 315-316 319
			329-331 345 359 384 417 423-424
adult kidney	Invitrogen	AKT002	3 6 14 20-21 25-26 76 79 85 89 94 101 111
,		1	114 118 121 124 126 130-131 138 146 163
'			170 177-178 189 196 198 201 204 213 231
	1		320 329 342
adult lung	GIBCO	ALG001	4 29 74 79 85 90 96 105 111 119 132 134
	·	· ·	136 142 144 149 159 181 189 198 200 205-
,			207 226 255 257 263 283 294 300 302-303
			328 358-359 365 426
lymph node	Clontech	ALNO01	6 16 31 105 120 215 257 295 306 309 359
young liver	GIBCO	ALV001	10-11 25-26 29 31 33 76 85 95 115 121-122
			124 126 130 143 146 156 158 164 178 182 187 189 229 248 253-254 261 278 283 304
			342 375
adult liver	Invitrogen	ALV002	10-12 23 26 31 33-34 38 53 56 90-92 94-95
			118 121 124 128-129 138 141 146 148 153
			156 161 171 178 198 216 232 248 253-254
			256-257 264 302 306 365 375 383 396
adult liver	Clontech	ALV003	10-11 156 171 188
Ovary	Invitrogen	AOV001	3-8 10-11 14 16 19-22 24 27-31 34 36 57 73
	ľ	·	75-76 81-82 85 89-91 94-98 104-109 111
			115-116 121-128 130-131 134 136 138-139
			141 143-144 146 149-150 152 155 157-160 163-166 170-173 175 177-178 180 182 184-
		1	187 189-190 193-194 196-197 200-201 212-
	1		213 215 217 222 225-226 228 230-233 235
			241-243 245 248 253-259 261 266-267 270
]	272-273 276-278 283-285 287 289 292 297-
	ļ		299 305-306 315-317 319 323-325 329-331
			341 343-344 352 358-359 363-366 382-383
73.	÷	222004	386 389-390 412
Placenta	Invitrogen	APL001	73 92 117 135 182 194 232 246 261 272 282 359
placenta	Invitrogen	APL002	16 28 92 121 135 144 157 178 210 394
adult spleen	GIBCO	ASP001	3-4 16 32-33 35 90 96 99-100 123-125 128
	·	}	131 134 136 139 151 178 181 189 194 200
			210 218 229 251 253-255 257 276 283 307-
<u> </u>			309 315 329 354-355 357 392 400
testis	GIBCO	ATS001	22 73 82 91 96-97 104-105 117 124 130 134
			164 173 200 209 222 233 241 253-254 257
hladdar.	Trusibuscan	Dr. D001	285 287-288 305 325 329 351-353 359
bladder	Invitrogen	BLD001	4 108 130 150 212 226 236 240 242 257 276 287 305 395-396 415
bone marrow	Clontech	BMD001	1 4-5 22 29-30 34 72 85 88 90 92 94 98
			104-107 109 111 113 117 120 123-125 128-
		1	129 132 135 140 142 144 146 152 163 165-
1	}	1	166 170-173 177 180 182 186 189-190 198-
			209 215 222 225 232 240-246 251-252 260-
		1	261 273-275 277-280 283-285 300 316 318
h ana	an	Dimono	346-347 359
bone marrow	GF	BMD002	1 4 7-8 10-11 16 19 25 31 49 61-62 72 74
			76 80 85 88 90 93-95 97-101 109-110 112
L		1	114 116-117 121 126 129 132 135 141 144

Table 1

Tissue Origin	RNA/Tissue	Library	SEQ ID NO:
TIDDEC OLLGEN	Source	Name	SEQ ID NO:
	BOULCE	Merme	146 149-150 154 157 160 162-163 165-166
	ł		170-172 175 178-180 182-183 186-190 192-
			194 198-200 203 208 210-213 215 223 225
			234 242 245 247 251-254 256-257 265 270
			273 276-278 280 285 287 289 291 293-294
			299 302 307 309 315 322 324 337-338 353
			356-357 359 367 369 388 407 414 419 426
	·		434
bone marrow	Clonetech	BMD007	144
*Mixture of	VARIOUS	CGd010	1 34-35 95 152 161 171 182 206 219 242 260
16 tissues -	VENDORS	000010	267 276 280 288 297 300 315-316 412
mRNA			
*Mixture of	Various	CGd011	45 51 167 188 216 251-252
16 tissues -	Vendors		
mRNA			·
*Mixture of	Various	CGd012	2 10-11 18-21 29 31 34-35 40 42-43 45 48
16 tissues -	Vendors		50-52 69-71 87-89 94-95 98-105 109 111-113
mRNA			117 120 123 125 127 131 135-136 138 146
			158 163 165-169 175 180 187-188 191 198
			201 208 216 219-221 224 226 234 236 238-
			239 241-246 251-252 260 264 270 276-277
İ			279 281 283-284 287 295-296 314 319 321
	•		327-328 331 333-334 337-341 343 351-352
			361 365 369 379-380 387 389 395 397-399
		ĺ	402 406 410-412 417 419 424 426 431-433
*Mixture of	Various	CGd013	29 48 101 146 167-169 187 219 234 327 333
16 tissues -	Vendors		339 341 365 412 433
mRNA			
*Mixture of	Various	CGd015	29 86 90 95 98 110 113 118 132 158 171 184
16 tissues -	Vendors		193 218-220 243 284 310 385 410 419
mRNA	<u> </u>		
*Mixture of	Various	CGd016	3-4 20-21 29 38 85 88-89 95 105 119 122
16 tissues -	Vendors		131-133 140 185 211-212 225 256-257 273
mRNA			276 302 318 379-380 390 400 419
colon	Invitrogen	CLN001	4 25 33 85 138 146 148 158-159 198 210 229
			301 360 384 397
cervix	BioChain	CVX001	3 5 10-11 18 20-21 24-25 29 36 41 47 57 63
	1		72 74 76 86 90 94 104 108-109 111 125 127
	1	1	130 134 138 144 147 162 174 178-179 182
		ļ	186 189 193 197 211 222 225-226 228 232
			241 243 257 261 267 270 273-275 278-281
			288-289 298 301-302 305 315 319 324-325
			329 331 337-338 359 391-392 395 420
endothelial	Strategene	EDT001	3-6 18-19 24 27-29 35 72 76 79-80 85 89 96
cells			98 104-107 111 117 119-121 124-131 134 136
1		ĺ	138-139 141 144 146-147 149 152 158-159
1	1		166-167 170-173 178-179 182-183 186-187
		<u> </u>	191 193-194 196-197 200 210-211 222-224
			226 231-232 236 241 243 246 248 253-256
	1	1	258-259 276 279 282 287 292 300 302-303
ļ	\	l	315 329 337-338 358-362 382-383 385-388
esophagus	BioChain	ESO002	257
fetal brain	Clontech	FBR001	34
fetal brain	Clontech	FBR004	3 139 144 271 284 337-338
fetal brain	Clontech	FBR006	4 6-11 14 18-21 24 28 31 37-38 40 63 76 85
L	<u> </u>	L	87 89-90 94-95 97 105 108-109 112-113 115

Table 1

Tissue Origin	RNA/Tissue	Library	SEQ ID NO:
	Sourch	Name	
			117-120 127-130 133 138 140 144-146 148
•	,	(170 172 175 180 182 186-188 190 192 194
			199 201 203 209-210 215 219 222 229-230
	i		232-233 240 243 245 253-255 270 273 276
		Ì	281 288-289 292 295 304 315 317 319 324
	{		330-331 356-357 359-360 364 367-368 379-
	{	1	380 383 389 397 399-401 408-409 411 413
			419 421 423
fetal brain	Invitrogen	FBT002	2 14 19 23 28 31 90 94 105 121 124 126 131
			135 139 142 149 158 186 193 198 210 214-
•	ŀ	1	215 232 239 242 248 255 267 326 332 365
	<u> </u>		369 371 376-383 394 399
fetal heart	Invitrogen	FHR001	4 7-8 10-11 14 17-21 28-29 31-32 60 64-65
		ļ	73 85 87 92 95 102-103 105 108 111 113 117
,			119 121 125 128-129 134-135 141 152 154
			156-157 160-161 172 176 178 194 196 198-
,	(l	200 203 208 212 215 218 222 226 229 233-
			234 253-257 261 265 272 276 281 292-293
			295 303 305 319 325 327 337-338 341 345
			417 436
fetal kidney	Clontech	FKD001	1 14 22 94 110 115 132 134-135 146 178 189
recar kruney	CTOHEECH	LYDOOT	199 235-236 242 247 257 267 292 295 359
fetal kidney	Clontech	FKD002	22 31 38 40 46 94 122 127 131 156 160 194
recar viole?	CIOncech	FREE	198 229 253-254 270 292 303 319 354-355
			389 396
fetal kidney	Invitrogen	FKD007	303
fetal lung	Clontech	FLG001	85 89 98-100 111 175 271 281 369
fetal lung	Invitrogen	FLG003	84 88 106-107 122 135 140 146 160 181 246
			272 284 292 328 330 396 404 416 426
fetal liver-	Soares	FLS001	1-3 6-12 14 19 23 28-31 33 57 59-60 72-76
spleen	ļ		78 80 83 85-138 140-141 143-144 146-155
<u> </u>	1.	ļ ·	157-161 163-197 200 204 208 210-211 223
,		ŀ	225 230 232-233 235 241-243 245-266 268-
	•	{	273 277 281 285-287 292 297 303 314 329
	<u> </u>		343 346-347 357-359 369 397 399 407 415
fetal liver-	Soares	FLS002	1 3-4 6 10-12 23-24 29 31-33 35-37 53-54
spleen	,	}	74-76 79 81-82 86-89 91 94-95 99-104 106-
	ļ	ł	109 111-112 115 117-120 122 125-126 128-
	[1	129 132 134 136-138 141 146 149 153 157-
,			159 162-166 170 172 175 178-180 183 185-
•	1	1	191 194 196-197 205 207-212 222-225 228
	İ		232-233 239-241 248 251-252 255-256 258-
Ì]		259 261-262 264 266-267 270-271 273-275
į ·	1	1	277-278 283 285 287 298 305 315 317-318
			322 330-332 337-338 341 343 349 357-360
Fatal 15	+ Connec	FF COO2	365 388 390-391 399 402 418 424
fetal liver-	Soares	FLS003	12 29 91 98 111 119 156 163 165 178 186 193 210-211 276 286 315 322 346-347 357
spleen	İ	1	193 210-211 276 286 315 322 346-347 357 365 424
Foto1 1:	Tresitore	ET YZO O 1	7-8 14 35 118 122-123 129 146 182 211 230
fetal liver	Invitrogen	FLV001	7-8 14 35 118 122-123 129 146 182 211 230
1			346-347 352 365 367-369
fetal liver	Clontech	FLV002	102-103 147 149 300
	Clontech	FLV002	73 85 105 108 118 122 126 141 156-157 161
fetal liver	CTOHCECH	LTAAAA#	165 170 178 180 182 194 215 218 225 240
L			TOO TIO TOO TOD TOA TOA STO STO STO STO

Table 1

Tissue Origin	RNA/Tissue	Library	SEQ ID NO:
	Source	Name	
·			242 247 251-252 292 330 337-338 369 407 411 440
fetal muscle	Invitrogen	FMS002	5 9 17-18 20-21 29 38 85 88 97 106-107 129
	4		131 136 150-152 155 165 170 179 182 192-
			193 212-213 229 234 242 258-259 270 282
			286 289 300 316 319 345 351 354-355 360
			389 396 408 410 437 439
fetal skin	Invitrogen	FSK001	2 4 7-8 29 33 42-43 49 51-52 58 74 82 85
			90 94 110-111 116 118 121 133 136 138-139
			145 151 154 156-157 161-162 172 181 184
			186 193 198 200 205 207 209-211 222 227-
			230 232 235 240 246 253-257 266 270 276
			292 295 299 316 318 323 330 332 337-340
			343 357 369 389 394-395 412 422 427
fetal skin	Invitrogen	FSK002	4 9 42 44 51 66 72 81 85 89-90 95 98 105
		ļ	112-114 119 121 129 133 135 162 172 179-
			182 197 200 208 210 231 243-244 272 304
			316 330 339 354-355 357 360 389 395 410
		ì	417 437
fetal spleen	BioChain	FSP001	157 223
umbilical	BioChain	FUC001	4-6 20-21 25 29 73-74 83 87 89-91 94 101
cord	_		109 120 123 125 128 130-131 133 141 143-
			144 147 149 154 161 165 173 175 179 184
			188 210-212 217 226 235 240 248 251-252
· · · · ·	•		257 262 267 270 277 293 305 307 316 319
		ļ	323 327 331 341 356 359 389 392 407 416
fetal brain	GIBCO	HFB001	2-4 16 20-21 74 77 85 89-91 96-98 104-105
·		i .	111 114 118 121-122 124-125 127-128 131
	•		134 137-140 142 144 146-148 151 153 158-
		1	159 163-164 166 173 178 180 182 191 194
			196 200 203 209-214 216-232 234-236 238-
			239 243 253-255 263 270 272-273 276 281
		l	292 310 316 319-321 332 348 357 359 365
·			399
macrophage	Invitrogen	HMP001	2 247
infant brain	Soares	IB2002	2-4 7-8 19-22 26-27 31-32 35 73-74 80 85
	[1	89 91 96-98 106-107 110 112 118-119 121-
			122 125 128-131 134-144 148 153 164 166
			172-173 177 180 186-187 191-194 196 202-
			203 208-210 217 219 223-224 227 229 232-
			234 236-237 239 241-243 245 248 253-259
	1		273-275 278-279 282 287 294 298 309 314
			317 322 327 330 333-334 341 348-350 360
		<u> </u>	368 376 379-380 382 396 406 424
infant brain	Soares	IB2003	3-4 20-21 26 28 31 35 73 85 95-96 110 113
			119 122-123 130-131 135 138 140 142-143
			146 153 155 170 172-173 186 191-193 196
			209 219 223 226 229 233-234 236 239 245
·			248 253-254 256-257 273 279 291-292 304
			314 337-338 343 359 367 371 376 397 413
lung,	Strategene	LFB001	3 6 31 72-73 90 92 105-107 124 126-127 133
fibroblast			136 139 144 146 172 189 198 204 233 235
· ·		1	246 258-259 268 272 276 282 310 335 359
			434
adult lung	Invitrogen	LGT002	4 19-21 28 33 35-36 49 72 79 81 85 88 90-
			91 94-95 101 106-107 109 118 120-125 127

Table 1

Minan - A-	I mare /	T = 33	T 222
Tissue Origin	RNA/Tissue	Library	SEQ ID NO:
<u> </u>	Source	Name	
	·	1	130-131 133 135-138 141-142 144 147 149
		1	157 159-161 163 166 170-173 193-194 196-
			197 212 216 218 221 223 226 228-229 231
	}		233 241 247-248 253-255 257 261 266-267
		1	270-275 277-278 282-283 292 298 301 303
	1	1	315 318 324 331 335 354-355 359 367 369
11	GTDGG		381 392-393 398
leukocytes	GIBCO	LUC001	1-5 15 19-21 28 30-33 37 72 74 91 94-95
			97-100 108-109 113 115 117 119-122 124-125
}		} .	127-128 134-138 141 144 146-148 150-151
·	}	}	157-158160 162-167 170-173 175-178 180-181
		Ì	187 189 192 194 197 200 212-213 215-216
		}	218-219 223 225 228-232 241-242 245-246
		į.	251-254 261 272-276 278-282 284 287-290
		<u> </u>	297-298 305 307 310-314 325 331 336 340
 	<u> </u>	<u> </u>	358-359 372 399 414
leukocytes	Clontech	LUC003	1 5 124 171 176 204 225 248 253-254 283
	<u> </u>		285 307 315
melanoma	Clontech	MEL004	4-5 24 37 72-74 81 85 106-107 113 136 177
	•	ł	203 205-207 209 231 243 284-285 315-316
	<u> </u>	<u> </u>	320 326 359 374 428
mammary gland	Invitrogen	MMG001	2 4-5 7-8 10-12 29 31 34-35 38 50 80-81 85
	1	1	89-90 92 94-97 105 108-109 119-124 126
	}		128-130 135 138-139 141-142 144 146-147
	<u> </u>	1	153 155 157-159 163 178-179 181-182 198
1	l	į	200 209-210 219 223 228 230 232-233 235-
i e	{ .	{	236 239 242 248 253-255 257 260-261 265-
	1	j	267 270 272 281 287 292 294 315-316 318
ĺ	{		324 327 330 337-340 354-355 357 369 372
	<u> </u>		383 392-395 401 404
neuron	Strategene	NTD001	35 47 89-90 111 118 164 232 253-254 276
			324 331 382
neuron	Strategene	NTR001	20-21 37 122 147-149 170 179 181 186 212
			226 258-259 265 276 369 436 438
neuronal	Strategene	NTU001	7-8 37 55 80 85 112 118 126-127 133 138
cells		}	140-141 151 170 181 210 214 225-226 236
	<u> </u>		243 287 328 330-331 357 383 400 436
pituitary	Clontech	PIT004	92 124 159 231
gland	L	<u> </u>	
placenta	Clontech	PLA003	34 46 88 126 128 159 182 186 197 201 267
[278 281-282 305 330 356 361 365 418
prostate	Clontech	PRT001	18 36 72 74 86 95 106-107 111 118 122 144
		<u> </u>	161 179 211 218 233 286 297
rectum	Invitrogen	REC001	9 31 85 121 128 147 171 200 219 257 292
			340 394 398 407 412
salivary	Clontech	SAL001	3 24 38 80 122 136 147 189 241 282 296 310
gland	1		351 392 395 415
saliva gland	Clontech	SALS03	118
small	Clontech	SIN001	12 16 25 82-83 89-90 93 95 98 105-109 111
intestine	{	1	122-123 125-128 133-134 137 139 142 161
	1		167 171 184 197 201 204 212 218 236 242-
1		1	243 248-249 253-254 257 267 276 284-285
1	}	1	292 297 300 303 310 313 317-318 325 340
į	!		343 352 354-355 359 383 391 416
spinal cord	Clontech	SPC001	3 39 84 86 94 96 105 115 117 130-131 134
-2			136 141 143 148 155 176 190-191 203 213
			,

Table 1

Tissue Origin	RNA/Tissue Source	Library Name	SEQ ID NO:
			224 233-234 236 239 279 283 298 320-321 332 336-338 356 359 365 404-406
thalamus	Clontech	THA002	2 20-21 23 74 81 85 105-106 116 121 131 146 171 185 188 200 209 219 233 239 256 258-259 273 276 362 399
thymus	Clonetech	THM001	16 29 33 57 80 82 85 90 93-94 106-107 120 126 128 134 141 161 176 194 223 228 235 253-254 261 274-275 278 285 298 319 332 336 343 353 359 425
thymus	Clontech	тнмс02	1-2 7-9 14 26 34 44 73 75 82 85 87 94 98 106-107 109-111 117 119-120 125-126 128- 129 139 141 144 147-148 151 154-155 162 165 170-172 175-176 179 182 186 193-194 199-200 208-209 213 218 233 235 240 242 247 253-254 257 265 276 281 287 290 305 307 312 319 336 342 354-356 359 364 367 399 408 412-413 415 419 421 426 429-433
thyroid gland	Clontech	THR001	3 5 7-8 28 30-31 33 73-77 80 82 85 88 90- 92 94 96-98 105-107 109 113 117 121-122 124-125 127-128 130 134 136 141 143 146- 148 152 161-163 166 175 177-178 181 194 199 201 204 210 212 216 218 223-226 228 230-231 234 236 241 243 246 253-257 261 270 272-273 276-278 281-283 287 292 295 298 303-304 308 315 323 329 335 352 359 362 401 416-417
trachea	Clontech	TRC001	88 138 180 226 228 279 359 411 436
uterus	Clontech	UTR001	3 10-11 23 77 92 106-107 109 111 141 197- 198 218 241 257 270 274-275 302 315 329 396 400 413

*The 16 tissue/mRNAs and their vendor sources are as follows: 1) Normal adult brain mRNA (Invitrogen), 2) Normal adult kidney mRNA (Invitrogen), 3) Normal fetal brain mRNA (Invitrogen), 4) Normal adult liver mRNA (Invitrogen), 5) Normal fetal kidney mRNA (Invitrogen), 6) Normal fetal liver mRNA (Invitrogen), 7) normal fetal skin mRNA (Invitrogen), 8) human adrenal gland mRNA (Clontech), 9) Human bone marrow mRNA (Clontech), 10) Human leukemia lymphoblastic mRNA (Clontech), 11) Human thymus mRNA (Clontech), 12) human lymph node mRNA (Clontech), 13) human so\spinal cord mRNA (Clontech), 14) human thyroid mRNA (Clontech), 15) human esophagus mRNA (BioChain), 16) human conceptional umbilical cord mRNA (BioChain).

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
1	gi9837125	Homo sapiens	membrane-associated nucleic acid binding protein mRNA, partial cds.	2553	54
1	gi7020305	Homo sapiens	cDNA FLJ20301 fis, clone HEP06569.	1728	47
1	gi7294120	Drosophila melanogaster	CG16807 gene product	1535	53
2	AAY57911	Homo sapiens	Human transmembrane protein HTMPN-35.	1258	82
2	AAB88406	Homo sapiens	Human membrane or secretory protein clone PSEC0162.	265	39
2	gi14272664	Homo sapiens	unnamed protein product	265	39
3	gi12654575	Homo sapiens	Similar to gp25L2 protein, clone MGC:2142 IMAGE:2967520, mRNA, complete cds.	1116	100
3	gi12845568	Mus musculus	putative	1099	98
3	gi996057	Homo sapiens	H.sapiens mRNA for gp25L2 protein.	1096	98
4	gi9971050	Homo sapiens	Human DNA sequence from clone RP11-526K24 on chromosome 20. Contains a novel gene, the 5' end of a novel gene, two CpG islands, ESTs, GSSs and STSs, complete sequence.	4348	99
4	AAB95086	Homo sapiens	Human protein sequence SEQ ID NO:16999.	3034	99
4	gi10433753	Homo sapiens	cDNA FLJ12307 fis, clone MAMMA1001908.	3034	99
5	gi4689106	Homo sapiens	NADH-ubiquinone oxidoreductase B8 subunit	505	100
5	gi2909862	Homo sapiens	NADH-ubiquinone oxidoreductase subunit CI-B8 mRNA, complete cds.	505	100
5	gi12539408	Homo sapiens	NDUFA2 gene for NADH dehydrogenase (ubiquinone) 1 alpha subcomplex 2, complete cds.	505	100
6	AAG64416	Homo sapiens	Human nucleoprotein.	3765	100
6	gi10443046	Homo sapiens	Human DNA sequence from clone RP11-465L10 on chromosome 20. Contains 10 CpG islands, ESTs, STSs and GSSs. Contains the gene for a novel protein similar to Drosophila CG11399, the gene for a novel C2H2 type zinc finger protein similar to chicken FZF-1, a Ferritin light polypeptide (FTL) pseudogene, the MMP9 gene for matrix metalloproteinase 9 (gelatinase B, 92kD gelatinase, 92kD type IV collagenase) (CLG4B), a novel gene, the SLC12A5 gene for solute carrier family 12, (potassium-chloride transporter) member 5 (KIAA1176) and the 3' end of gene KIAA1637, complete sequence.	3765	100
6	gi15426514	Homo sapiens	clone MGC:16205 IMAGE:3640928, mRNA, complete cds.	3765	100
7	AAG64416	Homo sapiens	Human nucleoprotein.	3366	100

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
7	gi10443046	Homo sapiens	Human DNA sequence from clone	3366	100
			RP11-465L10 on chromosome 20.		
			Contains 10 CpG islands, ESTs, STSs		j
	!		and GSSs. Contains the gene for a		1
			novel protein similar to Drosophila		1.
			CG11399, the gene for a novel C2H2		
			type zinc finger protein similar to		
			chicken FZF-1, a Ferritin light		ľ
	1		polypeptide (FTL) pseudogene, the	1	ł
			MMP9 gene for matrix		
			metalloproteinase 9 (gelatinase B,		
		·	92kD gelatinase, 92kD type IV	1	ļ
			collagenase) (CLG4B), a novel gene,	١.	
	İ	i	the SLC12A5 gene for solute carrier		
	ļ	:	family 12, (potassium-chloride		
	İ		transporter) member 5 (KIAA1176)		
	1		and the 3' end of gene KIAA1637, complete sequence.	· .	
7	gi15426514	Homo sapiens		2266	100
,	g115420514	nomo sapiens	clone MGC:16205 IMAGE:3640928,	3366	100
8	gi14571904	Rattus	mRNA, complete cds.	2146	0.5
	g1143/1904	norvegicus	lysosomal amino acid transporter 1	2145	85
8	AAE04910	Homo sapiens	Human transporter and ion channel-23	1220	-
•	AAEU491U	riomo sapiens		1239	56
8	gi7297404	Drosophila	(TRICH-23) protein.	027	42
	g1/29/404	melanogaster	CG13384 gene product	837	43
9	AAB73686	Homo sapiens	Human oxidoreductase protein ORP- 19.	1301	98
9 .	gi7291405	Drosophila melanogaster	T3dh gene product	808	59
9	gi5824752	Caenorhabditis	predicted using Genefinder-contains	685	52
		elegans	similarity to Pfam domain: PF00465	-	
•		ŭ	(Iron-containing alcohol		,
			dehydrogenases), Score=177.7, E-		
			value=1.9e-50, N=2~cDNA EST		
			EMBL:Z14517 comes from this gene;	}	
		· ·	cDNA EST yk18d4.3 comes from this		1
			gene~cDNA EST yk18d4.5 comes		
			from this gene; cDNA EST yk116f5.5	1	Ì
		,	comes from this gene~cDNA EST		
			yk132h3.3 comes from this gene;		
		,	cDNA EST yk73d10.3 comes from this]]. [*]
			gene~cDNA EST yk93e9.3 comes from		1
			this gene; cDNA EST yk132h3.5		
			comes from this gene~cDNA EST		
		,	yk73d10.5 comes from this gene;		
. 1		·	cDNA EST yk93e9.5 comes from this	1	
·			gene~cDNA EST yk135b6.5 comes]
* .	İ		from this gene; cDNA EST yk135b6.3		
			comes from this gene-cDNA EST		
			yk201e5.3 comes from this gene;		
,			cDNA EST yk268b1.3 comes from this		
			gene~cDNA EST yk261d6.3 comes		1

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	%
		·			Identity
			from this gene; cDNA EST yk262h11.3		
1		i '	comes from this gene~cDNA EST]	
			yk292h11.3 comes from this gene;		
		ł	cDNA EST yk304d8.3 comes from this	l	
			gene~cDNA EST yk344b7.3 comes		
			from this gene; cDNA EST yk351a6.3		
			comes from this gene~cDNA EST	:	
			yk366d9.3 comes from this gene;		İ
		•	cDNA EST yk368e3.3 comes from this	1	
•	,		gene~cDNA EST yk372c11.3 comes		
			from this gene; cDNA EST yk389g3.3	i	
	ľ		comes from this gene~cDNA EST	<u> </u>	
			yk422d2.3 comes from this gene;		
	-	• •	cDNA EST yk381d7.3 comes from this		
	{		gene~cDNA EST yk201e5.5 comes		
• '			from this gene; cDNA EST yk267f6.5		
			comes from this gene~cDNA EST	[
			yk268b1.5 comes from this gene;		
1	1		cDNA EST yk261d6.5 comes from this	i	
	·	'	gene~cDNA EST yk262h11.5 comes	ŀ	
			from this gene; cDNA EST yk292h11.5		
			comes from this gene~cDNA EST		
•	· ·		yk304d8.5 comes from this gene;		
1			cDNA EST yk344b7.5 comes from this		
			gene~cDNA EST yk368e3.5 comes		
	•		from this gene; cDNA EST yk372c11.5		
			comes from this gene~cDNA EST		
			yk351a6.5 comes from this gene;		
	· ·		cDNA EST yk366d9.5 comes from this		
			gene~cDNA EST yk389g3.5 comes		
i			from this gene; cDNA EST yk422d2.5		
			comes from this gene~cDNA EST		
			yk560f4.3 comes from this gene;		
			cDNA EST yk625h5.3 comes from this		
			gene~cDNA EST yk381d7.5 comes		
			from this gene; cDNA EST yk560f4.5		
			comes from this gene~cDNA EST		•
10	A A D72 606	77	yk625h5.5 comes from this gene	1.555	00
10	AAB73686	Homo sapiens	Human oxidoreductase protein ORP-	1552	99
10	-:7201405	D1"	19.	001	
10	gi7291405	Drosophila	T3dh gene product	891	56
10	1500 1500	melanogaster			
10	gi5824752	Caenorhabditis	predicted using Genefinder-contains	730	51
		elegans	similarity to Pfam domain: PF00465		
			(Iron-containing alcohol		
			dehydrogenases), Score=177.7, E-	, ;	
			value=1.9e-50, N=2~cDNA EST		
			EMBL:Z14517 comes from this gene;		
			cDNA EST yk18d4.3 comes from this		
			gene~cDNA EST yk18d4.5 comes		
			from this gene; cDNA EST yk116f5.5	'	
1			comes from this gene~cDNA EST		
	,		yk132h3.3 comes from this gene;		
			cDNA EST yk73d10.3 comes from this		
L	<u> </u>	<u> </u>			

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	%
<u> </u>					Identity
1			gene-cDNA EST yk93e9.3 comes from		
}	Ì		this gene; cDNA EST yk132h3.5		
	İ	1	comes from this gene-cDNA EST		,
		1	yk73d10.5 comes from this gene;		
		İ	cDNA EST yk93e9.5 comes from this		l
			gene~cDNA EST yk135b6.5 comes	ļ	ļ ·
	1		from this gene; cDNA EST yk135b6.3	ŀ	
[Į.	comes from this gene~cDNA EST	l	l
		1	yk201e5.3 comes from this gene;		İ
[1	1	cDNA EST yk268b1.3 comes from this	ł	1
İ			gene~cDNA EST yk261d6.3 comes	1	
		İ		1	1
1	1		from this gene; cDNA EST yk262h11.3	1	1
			comes from this gene-cDNA EST		ŀ
	1	1	yk292h11.3 comes from this gene;	•	Ì
·	İ]	cDNA EST yk304d8.3 comes from this		
			gene~cDNA EST yk344b7.3 comes	ļ	[
		1	from this gene; cDNA EST yk351a6.3		}
		1	comes from this gene~cDNA EST		۱.
			yk366d9.3 comes from this gene;	l	
		İ	cDNA EST yk368e3.3 comes from this	1	•
			gene~cDNA EST yk372c11.3 comes		
	j		from this gene; cDNA EST yk389g3.3	1	
,			comes from this gene~cDNA EST		
,	1	\	yk422d2.3 comes from this gene;		
	ļ ·		cDNA EST yk381d7.3 comes from this		
			gene~cDNA EST yk201e5.5 comes		
	·	·	from this gene; cDNA EST yk267f6.5		
	ļ		comes from this gene~cDNA EST		
			yk268b1.5 comes from this gene;		
	· .		cDNA EST yk261d6.5 comes from this		
,			gene-cDNA EST yk262h11.5 comes		
			from this gene; cDNA EST yk292h11.5		
	1		comes from this gene~cDNA EST		
				,	
			yk304d8.5 comes from this gene;		
		Ì	cDNA EST yk344b7.5 comes from this		
	l .		gene-cDNA EST yk368e3.5 comes		
	1		from this gene; cDNA EST yk372c11.5		·
-	1	[comes from this gene-cDNA EST		
		j	yk351a6.5 comes from this gene;		
			cDNA EST yk366d9.5 comes from this		
			gene~cDNA EST yk389g3.5 comes		
			from this gene; cDNA EST yk422d2.5		· .
			comes from this gene~cDNA EST		
	1		yk560f4.3 comes from this gene;		
			cDNA EST yk625h5.3 comes from this		
		1	gene~cDNA EST yk381d7.5 comes		
	1		from this gene; cDNA EST yk560f4.5		
•		·	comes from this gene-cDNA EST		
	1		yk625h5.5 comes from this gene		
11	AAB85166	Homo sapiens	Human Bcl-Gl polypeptide.	1598	87
11	gi14598300	Homo sapiens	unnamed protein product	1598	87
11	gi12584085	Homo sapiens	apoptosis regulator BCL-G long form	1598	87
	6112304003	Tromo sabiens		1230	0/
12	gi15077865	Mina managaritan	(BCLG) mRNA, complete cds.	1050	82
14	RIT201/902	Mus musculus	bullous pemphigoid antigen 1-b	1253	82

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
12	gi15077863	Mus musculus	bullous pemphigoid antigen 1-a	1253	82
12	gi6624582	Homo sapiens	Human DNA sequence from clone RP1-61B2 on chromosome 6p11.2-12.3 Contains isoforms 1 and 3 of BPAG1 (bullous pemphigoid antigen 1 (230/240kD), an exon of a gene similar to murine MACF cytoskeletal protein,	733	99
		•	STSs and GSSs, complete sequence.		
13	gi3702270	Homo sapiens	chromosome 19, cosmid R31408, complete sequence.	887	93
13	gi401845	Homo sapiens	ribosomal protein L18a mRNA, complete cds.	887	93
13	gi13960144	Homo sapiens	ribosomal protein L18a, clone MGC:4476 IMAGE:2961519, mRNA, complete cds.	887	93
14	AAB59090	Homo sapiens	Breast and ovarian cancer associated antigen protein sequence SEQ ID 798.	496	80
14	AAB44129	Homo sapiens	Human cancer associated protein sequence SEQ ID NO:1574.	453	81
14	gi14198321	Mus musculus	ribosomal protein L31	453	81
15	gi5689465	Homo sapiens	mRNA for KIAA1064 protein, partial cds.	5643	100
15	gi4884368	Homo sapiens	mRNA; cDNA DKFZp586L1220 (from clone DKFZp586L1220); partial cds.	1628	100
15	gi13161145	Homo sapiens	zinc finger protein mRNA, complete cds.	369	36
16	gi5870832	Mus musculus	skm-BOP1	2494	94
16	gi5870834	Mus musculus	skm-BOP2	2397	91
16	gi1809322	Mus musculus	t-BOP	2285	93
17	gi13938126	Mus musculus	RIKEN cDNA 3732409C05 gene	2678	98
17	gi12852375	Mus musculus	putative	2678	98
17	gi7024433	Torpedo marmorata	male sterility protein 2-like protein	2307	80
18	AAB95482	Homo sapiens	Human protein sequence SEQ ID NO:18007.	1572	67
18	gi14042809	Homo sapiens	cDNA FLJ14932 fis, clone PLACE1009639.	1572	67
18	gi12053165	Homo sapiens	mRNA; cDNA DKFZp434K0427 (from clone DKFZp434K0427); complete cds.	1572	67
19	gi7243159	Homo sapiens	mRNA for KIAA1389 protein, partial cds.	7842	99
19	gi4151328	Homo sapiens	high-risk human papilloma viruses E6 oncoproteins targeted protein E6TP1 alpha mRNA, complete cds.	3777	53
19	gi4151330	Homo sapiens	high-risk human papilloma viruses E6 oncoproteins targeted protein E6TP1 beta mRNA, complete cds.	3768	53
20	gi7243159	Homo sapiens	mRNA for KIAA1389 protein, partial cds.	7714	98
20	gi4151328	Homo sapiens	high-risk human papilloma viruses E6 oncoproteins targeted protein E6TP1	3806	54

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
			alpha mRNA, complete cds.		
20	gi4151330	Homo sapiens	high-risk human papilloma viruses E6 oncoproteins targeted protein E6TP1 beta mRNA, complete cds.	3797	53
21	AAB95328	Homo sapiens	Human protein sequence SEQ ID NO:17595.	753	61
21	AAB93757	Homo sapiens	Human protein sequence SEQ ID NO:13432.	753	61
21	AAB29657	Homo sapiens	Human membrane-associated protein HUMAP-14.	753	61
22	gi7673373	Homo sapiens	SCAN-related protein RAZ1 (RAZ1) mRNA, partial cds.	1104	100
22	AAG93274	Homo sapiens	Human protein HP10543.	900	100
22	AAB42846	Homo sapiens	Human ORFX ORF2610 polypeptide sequence SEQ ID NO:5220.	900	100
23	gi7242963	Homo sapiens	mRNA for KIAA1304 protein, partial cds.	5409	99
23	gi3413874	Homo sapiens	mRNA for KIAA0456 protein, partial cds.	3695	67
23	AAB30852	Homo sapiens	Amino acid sequence of human signal transduction protein SGT6-1.	3685	68
24	AAG64386	Homo sapiens	Human alcohol dehydrogenase 39.	1228	77
24	gi12861800	Mus musculus	putative	1083	66
24	gi3878713	Caenorhabditis elegans	weak similarity with quinone oxidoreductase, contains similarity to Pfam domain: PF00107 (Zinc-binding dehydrogenases), Score—80.6, E-value=6.2e-06, N=1~cDNA EST yk164b4.5 comes from this gene~cDNA EST yk164b4.3 comes from this gene~cDNA EST yk264f3.5 comes from this gene	556	39
25	AAE02629	Homo sapiens	Human secreted protein Zalpha37.	2481	100
25	gi14536691	Homo sapiens	unnamed protein product	2481	100
25	AAY99419	Homo sapiens	Human PRO1780 (UNQ842) amino acid sequence SEQ ID NO:282.	1960	77
26	gi6102869	Homo sapiens	mRNA; cDNA DKFZp434H1235 (from clone DKFZp434H1235); partial cds.	831	100
26	gi12853439	Mus musculus	putative	789	94
26	gi2198807	Gallus gallus	monocarboxylate transporter 3	505	29
27	gi7299069	Drosophila melanogaster	CG11755 gene product	205	34
27	gi3875367	Caenorhabditis elegans	contains 3 cysteine rich repeats	136	41
27	gi3249080	Arabidopsis thaliana	Contains similarity to MYB transcription factor isolog T01O24.1 gb 2288980 from A. thaliana BAC gb AC002335.	69	35
28	gi11041628	Homo sapiens	RPL6 gene for ribosomal protein L6, complete cds.	1207	98
28	gi433416	Homo sapiens	Human mRNA for DNA-binding protein, TAXREB107, complete cds.	1207	98

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
28	gi13278717	Homo sapiens	ribosomal protein L6, clone MGC:1635 IMAGE:2823733, mRNA, complete cds.	1207	98
29	AAG03810	Homo sapiens	Human secreted protein, SEQ ID NO: 7891.	845	100
29	gi186800	Homo sapiens	Human ribosomal protein L12 mRNA, complete cds.	845	100
29	gi14198333	Homo sapiens	ribosomal protein L12, clone MGC:9760 IMAGE:3855674, mRNA, complete cds.	845	100
30	AAB95051	Homo sapiens	Human protein sequence SEQ ID NO:16849.	2965	100
30	gi10433519	Homo sapiens	cDNA FLJ12118 fis, clone MAMMA1000085, weakly similar to PUTATIVE CYSTEINYL-TRNA SYNTHETASE C29E6.06C (EC 6.1.1.16).	2965	100
30	gi13938199	Homo sapiens	hypothetical protein FLJ12118, clone MGC:15044 IMAGE:2822557, mRNA, complete cds.	2959	99
31	gi12858123	Mus musculus	putative	2441	73
31	gi7959195	Homo sapiens	mRNA for KIAA1467 protein, partial cds.	2232	100
31	gi13278148	Mus musculus	Similar to RIKEN cDNA 8430419L09 gene	794	83
32	gi15530305	Homo sapiens	Similar to RIKEN cDNA 1700045119 gene, clone MGC:2647 IMAGE:3509621, mRNA, complete cds.	1245	84
32	gi9858803	Mus musculus	Zfp228	512	47
32	AAG75629	Homo sapiens	Human colon cancer antigen protein SEQ ID NO:6393.	511	46
33	gi8101071	Homo sapiens	golgin-like protein (GLP) gene, complete cds.	312	46
33	gi8099669	Homo sapiens	golgin-like protein (GLP) mRNA, complete cds.	312	46
33	gi11037008	Human herpesvirus 8	latent nuclear antigen	245	40
34	gi437985	Canis familiaris	Rab12 protein	1071	99
34	gi206531	Rattus norvegicus	RAB12	995	96
34	gi12851149	Mus musculus	putative	819	96
35	gi13543689	Homo sapiens	Similar to RIKEN cDNA 4933405K01 gene, clone MGC:14799 IMAGE:4068454, mRNA, complete cds.	1077	96
35	gi12805373	Mus musculus	Unknown (protein for MGC:7298)	950	84
35	gi12855529	Mus musculus	putative	642	79
36	gi12697979	Homo sapiens	mRNA for KIAA1717 protein, partial cds.	1982	100
36	gi1651678	Synechocystis sp. PCC 6803	ORF_ID:slr1485~hypothetical protein	185	34

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	%
36	gi2739367	Arabidopsis thaliana	putative phosphatidylinositol-4- phosphate 5-kinase	153	Identity 28
37	gi3800892	Homo sapiens	neurexin III-alpha gene, partial cds.	1255	99
37	gi294602	Rattus norvegicus	neurexin III-alpha	1160	91
37	gi205716	Rattus norvegicus	neurexin II-alpha-a	561	50
38	gi10047315	Homo sapiens	mRNA for KIAA1619 protein, partial cds.	4447	99
38	gi8217424	Homo sapiens	Human DNA sequence from clone RP11-108L7 on chromosome 10. contains part of the gene for a novel Insulin-like growth factor binding type protein with Kazal-type serine protease	4407	99
			inhibitor domain, the gene for a novel protein similar to rat tricarboxylate carrier, the gene for a novel PDZ (DHR, GLGF) domain protein, the gene for a novel protein similar to KIAA0552, KIAA0341 and Fugu		
			hypothetical protein 2, the gene for a novel protein similar to Plasmodium POM1 and C. elegans F46G11.1, a putative novel gene, the SEMA4G gene for semaphorin 4G and a novel gene. Contains ESTs, STSs, GSSs and seven		
			putative CpG islands, complete sequence.	-	
38	gi4836757	Mus musculus	semaphorin subclass 4 member G	4021	90
39	gi10438664	Homo sapiens	cDNA: FLJ22324 fis, clone HRC05551.	307	100
39	gi13559240	Homo sapiens	Human DNA sequence from clone RP5-842G6 on chromosome 20. Contains the 3' end of a novel gene, the 3' end of the gene for a novel protein similar to SEL1L (sel-1 (suppressor of lin-12, C.elegans)-like), ESTs, STSs and GSSs, complete sequence.	307	100
39	gi13543669	Homo sapiens	hypothetical protein FLJ22324, clone MGC:14701 IMAGE:4247211, mRNA, complete cds.	307	100
40	gi14595019	Homo sapiens	mRNA for keratin 6 irs (KRT6IRS gene).	2615	99
40	gi6092075	Mus musculus	type II cytokeratin	2414	01
10		Homo sapiens	Similar to keratin 6A, clone MGC:20671 IMAGE:3639270, mRNA, complete cds.	1468	91 57
11	gi12655452	Homo sapiens	mRNA for keratin associated protein 4.7 (KRTAP4.7 gene).	1157	86
1	gi12655464	Homo sapiens	partial mRNA for keratin associated protein 4.15 (KRTAP4.15 gene).	1090	88
1	gi12655462	Homo sapiens	mRNA for keratin associated protein 4.14 (KRTAP4.14 gene).	1063	84

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
42	gi553772	Homo sapiens	Human Tcr-C-delta gene, exons 1-4; Tcr-V-delta gene, exons 1-2; T-cell receptor alpha (Tcr-alpha) gene, J1-J61 segments; and Tcr-C-alpha gene, exons 1-4.	110	100
42	gi4379087	Homo sapiens	mRNA for TCR alpha variable region, patient AF31.	73	46
42	AAW40057	Homo sapiens	Cellular transcriptional factor p300.	71	42
43	gi15866589	Capsella rubella	hypothetical protein	97	30
43	gi3879045	Caenorhabditis elegans	R102.6	96	34
43	AAY56133	Homo sapiens	Human N-methyl-D-aspartate receptor 2 subunit SEQ ID NO:54.	94	52
44	gi13569345	Homo sapiens	pregnancy-associated plasma preproprotein-A2 mRNA, complete cds.	9839	99
44	gi10639043	Homo sapiens	mRNA for pregnancy-associated plasma protein-E (PAPPE gene).	8966	99
44	gi1142970	Homo sapiens	Human pregnancy-associated plasma protein-A preproform (PAPPA) mRNA, complete cds.	3856	45
45	gi12851017	Mus musculus	putative	578	83
45	gi4490653	Schizosacchar omyces pombe	profilin.	186	35
45	gi440266	Acanthamoeba castellanii	profilin I	166	34
46	gi1617480	Comamonas testosteroni	unknown	712	82
46	gi3046394	Ralstonia eutropha	phbF	563	66
46	gi6683782	Burkholderia sp. DSMZ 9242	unknown	560	61
47	gi9229934	Mus musculus	midnolin	2103	78
47	AAB56832	Homo sapiens	Human prostate cancer antigen protein sequence SEQ ID NO:1410.	912	71
47	gi15929300	Homo sapiens	Similar to midnolin, clone IMAGE:3958934, mRNA, partial cds.	907	100
48	gi13377624	Homo sapiens	calicin mRNA, complete cds.	3089	99
48	gi854100	Homo sapiens	H.sapiens mRNA for calicin (partial).	3076	99
48	gi853784	Bos taurus	calicin	2896	91
49	AAB68411	Homo sapiens	Amino acid sequence of a human NOV2 polypeptide.	2131	100
49	AAY99407	Homo sapiens	Human PRO1337 (UNQ692) amino acid sequence SEQ ID NO:236.	2101	99
49	AAB68414	Homo sapiens	Amino acid sequence of NOV2 polypeptide clone TA-cgAL132708 A.	2014	99
50	gi12082748	Mus musculus	T-box transcription factor TBX18	2972	93
50	gi5102617	Homo sapiens	Human DNA sequence from clone 33L1 on chromosome 6q14.1-15. Contains the gene for novel T-box (Brachyury) family protein. Contains	2634	100

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
			ESTs, STSs, GSSs and two putative CpG islands, complete sequence.		
50	gi12849661	Mus musculus	putative	2223	96
51	gi12843048	Mus musculus	putative	339	72
51	gi6691626	Homo sapiens	RAGE mRNA for advanced glycation endproducts receptor, complete cds.	111	32
51	gi190846	Homo sapiens	Human receptor for advanced glycosylation end products (RAGE) mRNA, partial cds.	111	32
52	AAG71840	Homo sapiens	Human olfactory receptor polypeptide, SEQ ID NO: 1521.	1313	85
52	AAG71839	Homo sapiens	Human olfactory receptor polypeptide, SEQ ID NO: 1520.	1226	81
52	AAG71837	Homo sapiens	Human olfactory receptor polypeptide, SEQ ID NO: 1518.	1159	-77
53	AAB94026	Homo sapiens	Human protein sequence SEQ ID NO:14163.	966	98
53	gi10433955	Homo sapiens	cDNA FLJ12457 fis, clone NT2RM1000666, weakly similar to DNA-BINDING PROTEIN A.	966	98
53	gi7295442	Drosophila melanogaster	CG17334 gene product	302	47
54	gi8980396	Homo sapiens	mRNA for T-cell antigen receptor- alpha, clone Pil-1a, partial.	566	97
54	gi2358063	Homo sapiens	T-cell receptor alpha delta locus from bases 752679 to 1000555 (section 4 of 5) of the Complete Nucleotide Sequence.	565	100
54	gi623149	Macaca mulatta	T-cell receptor alpha	512	85
55	gi2792496	Rattus norvegicus	tulip 2	2437	86
55	gi4884288	Homo sapiens	mRNA; cDNA DKFZp566D133 (from clone DKFZp566D133); partial cds.	1983	99
55	AAB41763	Homo sapiens	Human ORFX ORF1527 polypeptide sequence SEQ ID NO:3054.	1976	98
56	gi15524592	Homo sapiens	unnamed protein product	1033	52
56	gi537514	Homo sapiens	Human arylacetamide deacetylase mRNA, complete cds.	1033	52
56	AAB54079	Homo sapiens	Human pancreatic cancer antigen protein sequence SEQ ID NO:531.	1017	51
57	AAB33831	Homo sapiens	Human secreted protein BLAST search protein SEQ ID NO: 175.	149	35
57	gi1109682	Bos taurus	G-protein gamma-12 subunit	149	35
57	AAW09416	Homo sapiens	Human G protein gamma-7 subunit.	144	33
58	gi12082750	Mus musculus	T-box transcription factor TBX20	1469	93
58	gi9909810	Mus musculus	T-box transcription factor	1469	93
58	gi7229717	Danio rerio	H15-related T-box transcription factor hrT	1346	85
59	gi4185946	Human endogenous retrovirus K	gag protein	146	26
59	gi5802821	Homo sapiens	endogenous retrovirus HERV-K108,	146	26

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
_			complete sequence.		
59	gi5802814	Homo sapiens	endogenous retrovirus HERV-K103, complete sequence.	146	26
60	AAB94756	Homo sapiens	Human protein sequence SEQ ID NO:15815.	126	42
60	gi332612	Gibbon ape leukemia virus	pol polyprotein	113	50
60	gi3133302	Sus scrofa	pol protein	110	53
61	gi10121625	Gillichthys mirabilis	60S acidic ribosomal protein P1	127	81
61	AAB44012	Homo sapiens	Human cancer associated protein sequence SEQ ID NO:1457.	125	78
61	AAB43434	Homo sapiens	Human cancer associated protein sequence SEQ ID NO:879.	125	78
62	AAB12585	Homo sapiens	Human T cell activating protein SEQ ID NO:4.	140	37
62	gi12805221	Mus musculus	lymphocyte antigen 6 complex	140	37
62	gi198924	Mus musculus	Ly-6A.2	140	37
63	gi6969165	Homo sapiens	Human DNA sequence from clone RP3-475N16 on chromosome 6p12.3- 21.2. Contains the genes for CTG4A,	573	67
			pre-T cell receptor alpha, a novel protein similar to RPL7A (60S ribosomal protein L7A) and the 3' end of gene KIAA0240. Contains ESTs, STSs, GSSs and four putative CpG		
	1		islands, complete sequence.	1	
63	gi12841727	Mus musculus	putative	512	59
63	gi15293877	Ictalurus punctatus	ribosomal protein L7	314	38
64	gi181573	Homo sapiens	Human cytokeratin 8 (CK8) gene, complete cds.	1147	79
64	gi181400	Homo sapiens	Human cytokeratin 8 mRNA, complete cds.	1147	78
64	gi400416	Homo sapiens	H.sapiens KRT8 mRNA for keratin 8.	1147	79
65	gi13620887	Mus musculus	mitochondrial ribosomal protein S6	633	100
65	gi13620885	Homo sapiens	MRPS6 mRNA for mitochondrial ribosomal protein S6, partial cds.	565	85
65	gi14603226	Homo sapiens	clone MGC:19576 IMAGE:4304420, mRNA, complete cds.	565	85
66	gi13537119	Homo sapiens	mRNA for PAR-6 gamma, complete cds.	1956	100
66	gi8037909	Mus musculus	PAR6A	1490	76
66	gi9453884	Homo sapiens	mRNA for 16-5-5, partial cds.	1304	93
67	AAB95293	Homo sapiens	Human protein sequence SEQ ID NO:17517.	776	79
67	AAG81270	Homo sapiens	Human AFP protein sequence SEQ ID NO:58.	776	79
67	gi14035848	Homo sapiens	unnamed protein product	776	79
68	gi7020759	Homo sapiens	cDNA FLJ20565 fis, clone REC00542.	930	60
68	gi15216181	Homo sapiens	mRNA for putative 67-11-3 protein.	927	60
68	gi15930069	Homo sapiens	Similar to hypothetical protein FLJ20565, clone MGC:8850	917	60

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
			IMAGE:3914396, mRNA, complete cds.		
69	gi3228237	Homo sapiens	UHS KerB gene.	810	72
69	gi200962	Mus musculus	serine 1 ultra high sulfur protein	755	69
69	gi32472	Homo sapiens	H.sapiens mRNA for high-sulphur keratin.	749	71
70	AAB92789	Homo sapiens	Human protein sequence SEQ ID NO:11284.	3518	100
70	gi7022420	Homo sapiens	cDNA FLJ10407 fis, clone NT2RM4000520.	3518	100
70	gi13111786	Homo sapiens	hypothetical protein FLJ10407, clone MGC:970 IMAGE:3509727, mRNA, complete cds.	3511	99
71	gi13325178	Homo sapiens	Similar to RIKEN cDNA 2210016F16 gene, clone MGC:10999 IMAGE:3638524, mRNA, complete cds.	856	100
71	gi7291278	Drosophila melanogaster	CG9752 gene product	744	43
71	gi2854153	Caenorhabditis elegans	Hypothetical protein C11D2.4	729	45
72	gi7020991	Homo sapiens	cDNA FLJ20718 fis, clone HEP17872.	3013	100
72	gi15680144	Homo sapiens	hypothetical protein FLJ20718, clone IMAGE:4577269, mRNA, partial cds.	2906	99
72	gi10801646	Macaca fascicularis	hypothetical protein	1097	99
73	AAG93290	Homo sapiens	Human protein HP10650.	1215	100
73	gi14587195	Homo sapiens	FAPP1-associated protein 1 (FASP1) mRNA, complete cds.	1215	100
73	gi8118225	Homo sapiens	chromosome 21 unknown mRNA.	1215	100
74	gi10436998	Homo sapiens	cDNA: FLJ21011 fis, clone CAE04289.	2522	100
74	gi15030282	Homo sapiens	clone MGC:16827 IMAGE:3855873, mRNA, complete cds.	2522	100
74	gi8570641	Homo sapiens	clone 133K02 unknown mRNA.	2514	99
75	gi6599255	Homo sapiens	mRNA; cDNA DKFZp434C0328 (from clone DKFZp434C0328).	1612	100
75	gi6330416	Homo sapiens	mRNA for KIAA1201 protein, partial cds.	554	38
75	AAB74726	Homo sapiens	Human membrane associated protein MEMAP-32.	496	35
76	gi7021059	Homo sapiens	cDNA FLJ20758 fis, clone HEP01508.	1450	100
76	AAW88552	Homo sapiens	Secreted protein encoded by gene 19 clone HSAVU34.	1429	100
76	gi15341707	Homo sapiens	clone MGC:19979 IMAGE:3939273, mRNA, complete cds.	1429	100
77	AAB95410	Homo sapiens	Human protein sequence SEQ ID NO:17796.	774	100
77	gi10435394	Homo sapiens	cDNA FLJ13391 fis, clone PLACE1001241.	774	100
77	gi10503974	Homo sapiens	clone SP24 unknown mRNA.	765	99
78	gi7020587	Homo sapiens	cDNA FLJ20467 fis, clone KAT06638.	737	100

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
78	AAB42883	Homo sapiens	Human ORFX ORF2647 polypeptide sequence SEQ ID NO:5294.	530	100
78	AAB56642	Homo sapiens	Human prostate cancer antigen protein sequence SEQ ID NO:1220.	530	100
79	AAW93948	Homo sapiens	Human regulatory molecule HRM-4 protein.	441	91
79	gi12852696	Mus musculus	putative	386	47
79	gi12751103	Homo sapiens	PNAS-129 mRNA, complete cds.	348	100
80	gi7243053	Homo sapiens	mRNA for KIAA1336 protein, partial cds.	3851	99
80	gi7292144	Drosophila melanogaster	CG2069 gene product	1634	44
80	gi1065457	Caenorhabditis elegans	C54G7.4 gene product	706	25
81	gi10439581	Homo sapiens	cDNA: FLJ23023 fis, clone LNG01678.	652	100
81	gi7021132	Homo sapiens	cDNA FLJ20813 fis, clone ADSE01247.	652	100
81	AAG74674	Homo sapiens	Human colon cancer antigen protein SEQ ID NO:5438.	556	92
82	gi5262611	Homo sapiens	mRNA; cDNA DKFZp434I114 (from clone DKFZp434I114); complete cds.	838	100
82	gi11493368	Homo sapiens	Human DNA sequence from clone RP5-1009E24 on chromosome 20 Contains the SN gene encoding sialoadhesin, a novel gene similar to KIAA0417, the CENPB gene for centromere protein B, the CDC25B gene for Cell division cycle protein 25B, three novel genes, the 5' end of gene KIAA1271, nine CpG islands, ESTs, STSs and GSSs, complete	838	100
			sequence.	ļ. <u></u>	
82	gi13543798	Mus musculus	RIKEN cDNA 4931426K16 gene	680	92
83	AAB57003	Homo sapiens	Human prostate cancer antigen protein sequence SEQ ID NO:1581.	1302	99
83	AAR60558	Homo sapiens	Human basigin I.	1302	99
83	gi3492872	Homo sapiens	chromosome 19, cosmid F18382 (LLNLF-140D2) and 3' overlapping restriction fragment, complete sequence.	1302	99
84	gi9187614	Homo sapiens	mRNA full length insert cDNA clone EUROIMAGE 1759349.	580	100
84	AAB01394	Homo sapiens	Neuron-associated protein.	70	39
84	AAB54358	Homo sapiens	Human pancreatic cancer antigen protein sequence SEQ ID NO:810.	70	39
85	gi15986445	Homo sapiens	p90 autoantigen mRNA, complete cds.	4513	99
85	gi7959315	Homo sapiens	mRNA for KIAA1524 protein, partial cds.	4357	99
85	AAB95207	Homo sapiens	Human protein sequence SEQ ID NO:17311.	2341	100
86	gi7959231	Homo sapiens	mRNA for KIAA1485 protein, partial cds.	5813	99

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
86	AAB40418	Homo sapiens	Human ORFX ORF182 polypeptide sequence SEQ ID NO:364.	708	99
86	gi5901529	Homo sapiens	C2H2 type Kruppel-like zinc finger protein splice variant b (ZNF236) mRNA, complete cds.	520	24
87	gi7243270	Homo sapiens	mRNA for KIAA1436 protein, partial cds.	4604	99
87	gi5051974	Mus musculus	F2 alpha prostoglandin regulatory protein	4195	89
87	gì1054884	Rattus norvegicus	prostaglandin F2a receptor regulatory protein precursor	4191	88
88	gi13241286	Mus musculus	GABA(A) receptor-associated protein- like 2	607.	100
88	gi2104570	Rattus norvegicus	GEF-2	607	100
88	gi4433387	Bos taurus	general protein transport factor p16	607	100
89	gi15859535	Homo sapiens	unnamed protein product	5935	99
89	gi3043606	Homo sapiens	mRNA for KIAA0541 protein, partial cds.	5890	100
89	gi15624075	Homo sapiens	TGF-beta resistance-associated protein TRAG (TRAG) mRNA, partial cds.	5719	96
90	gi337370	Homo sapiens	Human rapamycin- and FK506-binding protein, complete cds.	740	100
90	gi13097252	Homo sapiens	Similar to FK506 binding protein 2 (13 kDa), clone MGC:5177 IMAGE:3445148, mRNA, complete cds.	740	100
90	AAQ31004_aa	Homo sapiens	hRFKBP cDNA.	735	99
91	gi12053147	Homo sapiens	mRNA; cDNA DKFZp434F1726 (from clone DKFZp434F1726).	1450	100
91	gi412195	Homo sapiens	unknown	265	98
91	AAR04931	Homo sapiens	Interferon-gamma receptor segment from clone 39 responsible for binding the target.	260	96
92	gi10437948	Homo sapiens	cDNA: FLJ21783 fis, clone HEP00284.	3276	100
92	AAB95352	Homo sapiens	Human protein sequence SEQ ID NO:17643.	1953	99
92	gi10435077	Homo sapiens	cDNA FLJ13171 fis, clone NT2RP3003819.	1953	99
93	gi12803319	Homo sapiens	clone MGC:3090 IMAGE:3347913, mRNA, complete cds.	4837	99
93	gi14044064	Homo sapiens	hypothetical protein DKFZp762M115, clone MGC:14418 IMAGE:4302613, mRNA, complete cds.	4831	99
93	gi10047337	Homo sapiens	mRNA for KIAA1630 protein, partial cds.	4671	100
94	AAB70535	Homo sapiens	Human PRO5 protein sequence SEQ ID NO:10.	2979	100
94	gi13185719	Homo sapiens	unnamed protein product	2979	100
94	AAB94106	Homo sapiens	Human protein sequence SEQ ID NO:14334.	2334	100
95	gi12837873	Mus musculus	putative	2370	75

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Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
95	gi13195574	Mus musculus	Praja1 isoform a	2339	75
95	AAB93847	Homo sapiens	Human protein sequence SEQ ID NO:13691.	1941	99
96	gi2224543	Homo sapiens	Human mRNA for KIAA0301 gene, partial cds.	10626	100
96	gi7529572	Homo sapiens	Human DNA sequence from clone RP1-122O8 on chromosome 6q14.2- 16.1. Contains the 3' part of a novel gene partially coded for by KIAA0301, a novel gene and the 3' part of the gene KIAA0957. Contains ESTs, STSs, GSSs and a putative CpG island, complete sequence.	10626	100
96	gi10727627	Drosophila melanogaster	CG13185 gene product	1452	34
97	AAB82318	Homo sapiens	Human immunoglobulin receptor IRTA5 protein.	2235	100
97	gi15528831	Homo sapiens	Fc receptor-like protein 1 (FCRH1) mRNA, complete cds.	2235	100
97	gi9930921	Homo sapiens	Human DNA sequence from clone RP11-367J7 on chromosome 1. Contains (part of) two or more genes for novel Immunoglobulin domains containing proteins, a SON DNA binding protein (SON) pseudogene, a voltage-dependent anion channel 1 (VDAC1) (plasmalemmal porin) pseudogene, ESTs, STSs and GSSs, complete sequence.	1533	100
98	AAB82318	Homo sapiens	Human immunoglobulin receptor IRTA5 protein.	2177	98
98	gi15528831	Homo sapiens	Fc receptor-like protein 1 (FCRH1) mRNA, complete cds.	2177	98
98	gi9930921	Homo sapiens	Human DNA sequence from clone RP11-367J7 on chromosome 1. Contains (part of) two or more genes for novel Immunoglobulin domains containing proteins, a SON DNA binding protein (SON) pseudogene, a voltage-dependent anion channel 1 (VDAC1) (plasmalemmal porin) pseudogene, ESTs, STSs and GSSs, complete sequence.	1533	100
99	gi10438861	Homo sapiens	cDNA: FLJ22461 fis, clone HRC10107.	4904	100
99	gi15079400	Homo sapiens	clone MGC:16796 IMAGE:3855477, mRNA, complete cds.	4899	99
99	AAU03497	Homo sapiens	Human sterol sensing domain protein.	4047	99
100	gi6524024	Mus musculus	mammalian inositol hexakisphosphate kinase 1	1031	50
100	gi10280996	Rattus norvegicus	inositol hexakisphosphate kinase	1027	49
100	gi6683115	Homo sapiens	mRNA for KIAA0263 protein, partial	1021	49

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
	†		cds.	 	lucuaty
101	gi6524024	Mus musculus	mammalian inositol hexakisphosphate kinase 1	1037	51
101	gi10280996	Rattus norvegicus	inositol hexakisphosphate kinase	1033	50
101	gi6683115	Homo sapiens	mRNA for KIAA0263 protein, partial cds.	1027	-50
102	gi13623311	Homo sapiens	clone IMAGE:3948563, mRNA, partial cds.	1629	100
102	gi3135968	Homo sapiens	Human DNA sequence from clone XXbac-3418 on chromosome 6p21.3- 22.1. Contains the 5' end of the ZNF184 gene for Kruppel-like zinc finger protein 184, a heterogeneous nuclear ribonucleoprotein A1	1627	47
			(HNRPA1) pseudogene, a CD83 antigen pseudogene, ESTs, STSs, GSSs and three CpG islands, complete sequence.		
102	gi1769491	Homo sapiens	Human kruppel-related zinc finger protein (ZNF184) mRNA, partial cds.	1625	47
103	gi16198398	Homo sapiens	clone MGC:27353 IMAGE:4671816, mRNA, complete cds.	2606	85
103	gi829151	Homo sapiens	H.sapiens ZNF37A mRNA for zinc finger protein.	1371	99
103	gi9801232	Homo sapiens	Human DNA sequence from clone RP11-508N22 on chromosome 10 Contains part of a novel gene (HSPC025), part of the ZNF37A (zinc finger protein 37a (KOX 21)) gene, part of a putative novel gene, ESTs, STSs, GSSs and a CpG Island, complete sequence.	1371	99
104	gi12053123	Homo sapiens	mRNA; cDNA DKFZp434K1421 (from clone DKFZp434K1421); complete cds.	2624	100
104	gi7292866	Drosophila melanogaster	CG15747 gene product	362	31
104	gi7549210	Babesia bigemina	200 kDa antigen p200	298	21
105	gi12053123	Homo sapiens	mRNA; cDNA DKFZp434K1421 (from clone DKFZp434K1421); complete cds.	2898	100
105	gi6841130	Homo sapiens	HSPC095 mRNA, partial cds.	419	100
105	gi7292866	Drosophila melanogaster	CG15747 gene product	364	30
106	gi10438207	Homo sapiens	cDNA: FLJ21977 fis, clone HEP05976.	1978	99
106	gi15012167	Homo sapiens	hypothetical protein FLJ21977, clone MGC:14918 IMAGE:3936410, mRNA, complete cds.	1974	99
106	AAB42499	Homo sapiens	Human ORFX ORF2263 polypeptide sequence SEQ ID NO:4526.	1392	100
107	gi1228035	Homo sapiens	Human mRNA for KIAA0191 gene,	8020	99

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
			partial cds.		
107	gi12697967	Homo sapiens	mRNA for KIAA1711 protein, partial cds.	1593	58
107	AAB94636	Homo sapiens	Human protein sequence SEQ ID NO:15515.	1004	52
108	AAG81252	Homo sapiens	Human AFP protein sequence SEQ ID NO:22.	2146	99
108	gi14035812	Homo sapiens	unnamed protein product	2146	99
108	gi10440123	Homo sapiens	cDNA: FLJ23436 fis, clone HRC12692.	2054	100
109	gi200009	Mus musculus	myosin I	5386	96
109	gi1666471	Mus musculus	myosin I heavy chain	5360	94
109	gi56733	Rattus norvegicus	myosin I heavy chain	5268	91
110	gi12053045	Homo sapiens	mRNA; cDNA DKFZp434K1115 (from clone DKFZp434K1115); complete cds.	4840	100
110	AAB65631	Homo sapiens	Novel protein kinase, SEQ ID NO: 158.	4835	99
110	gi14133215	Homo sapiens	mRNA for KIAA0781 protein, partial cds.	4678	100
111	gi12642596	Homo sapiens	nuclear receptor co-repressor/HDAC3 complex subunit TBLR1 (TBLR1) mRNA, complete cds.	2725	100
111	AAB95225	Homo sapiens	Human protein sequence SEQ ID NO:17352.	2720	99
111	gi10434648	Homo sapiens	cDNA FLJ12894 fis, clone NT2RP2004170, moderately similar to Homo sapiens mRNA for transducin (beta) like 1 protein.	2720	99
112	gi2224557	Homo sapiens	Human mRNA for KIAA0308 gene, partial cds.	6666	99
112	AAY23330	Homo sapiens	Human tumour suppressor (kismet) protein.	5759	98
112	gi7243213	Homo sapiens	mRNA for KIAA1416 protein, partial cds.	5264	59
113	gi12856019	Mus musculus	putative	1527	95
113	gi3947604	Caenorhabditis elegans	cDNA EST yk129f1.3 comes from this gene~cDNA EST yk129f1.5 comes from this gene~cDNA EST yk203e4.3 comes from this gene~cDNA EST yk191a9.3 comes from this gene~cDNA EST yk262c10.3 comes from this gene~cDNA EST yk278f9.3 comes from this gene~cDNA EST yk278f9.3 comes from this gene~cDNA EST yk325c7.3 comes from this gene~cDNA EST yk449a2.3 comes from this gene~cDNA EST yk449a2.3 comes from this gene~cDNA EST yk203e4.5 comes from this	787	41
			gene-cDNA EST yk191a9.5 comes from this gene-cDNA EST yk278f9.5 comes from this gene-cDNA EST yk262c10.5 comes from this		

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
			gene~cDNA EST yk325c7.5 comes		Auenuty
			from this gene~cDNA EST yk337f1.5		
		1	comes from this gene~cDNA EST		
		J	yk448g10.5 comes from this		1
		l	gene~cDNA EST yk449a2.5 comes	į	Į
			from this gene~cDNA EST yk636e2.3		
	i	İ	comes from this gene~cDNA EST	1	
			yk636e2.5 comes from this	ł	1
	1		gene~cDNA EST yk550e8.3 comes		
•	İ		from this gene~cDNA EST yk557a9.3		
			comes from this gene~cDNA EST	İ	
]		yk579c12.3 comes from this]
		,	gene~cDNA EST yk614e7.3 comes	1	
			from this gene-cDNA EST yk653f1.3	1	
		Į	comes from this gene~cDNA EST	1	
]	yk672b2.3 comes from this	1	
			gene~cDNA EST yk550e8.5 comes		
	}		from this gene~cDNA EST yk556b1.5		•
			comes from this gene~cDNA EST		
			yk557a9.5 comes from this		
	· ·		gene~cDNA EST yk579c12.5 comes		
			from this gene~cDNA EST yk606c8.5	1	
•			comes from this gene-cDNA EST	1	
	·		yk614e7.5 comes from this gene		
113	gi3947603	Caenorhabditis	cDNA EST yk167h7.3 comes from this	787	41
	J	elegans	gene~cDNA EST yk167h7.5 comes	107	47
			from this gene~cDNA EST yk289g5.3	ŀ	
			comes from this gene-cDNA EST	i	
			yk332h9.3 comes from this		
	•.		gene~cDNA EST yk289g5.5 comes		
			from this gene~cDNA EST yk332h9.5		
٠			comes from this gene~cDNA EST		
•			yk391h4.5 comes from this		
			gene~cDNA EST yk653f1.5 comes	\	
114	~:0200126	M	from this gene		
114	gi9280136	Macaca	unnamed protein product	3431	95
14	~:4060617	fascicularis			
114	gi4262617	Caenorhabditis	contains similarity to dual specificity	470	35
		elegans	phosphatase, catalyitic domain		
•		•	(Pfam:PF00782, Score=16.8, E=7.4e-		
	1000 (000		05, N=1)		
114	gi5706724	Homo sapiens	Cdc14B3 phosphatase mRNA,	166	30
			complete cds.		
115	AAB95254	Homo sapiens	Human protein sequence SEQ ID	3114	99
		· · · · · · · · · · · · · · · · · · ·	NO:17423.		
115	gi14042385	Homo sapiens	cDNA FLJ14693 fis, clone	3114	99 .
Ì			NT2RP2005360, weakly similar to		
	· .		Homo sapiens sentrin/SUMO-specific		
			protease (SENP1) mRNA.		
115	gi10314023	Homo sapiens	sentrin-specific protease (SENP2)	3107	99
	-		mRNA, complete cds.		
116	gi4240227	Homo sapiens	mRNA for KIAA0869 protein, partial	4417	98
			cds.	174/	70

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
116	gi13879506	Mus musculus	Unknown (protein for IMAGE:3963643)	4063	89
116	AAB93267	Homo sapiens	Human protein sequence SEQ ID NO:12300.	1895	97
117	gi13235092	Homo sapiens	mRNA for testis specific protein A14 (TSGA14 gene).	1957	100
117	gi10438839	Homo sapiens	cDNA: FLJ22445 fis, clone HRC09438.	1950	99
117	gi13235344	Mus musculus	testis specific protein a14	1704	87
118	gi7959279	Homo sapiens	mRNA for KIAA1509 protein, partial cds.	6769	99
118	AAB94101	Homo sapiens	Human protein sequence SEQ ID NO:14322.	1871	99
118	gi10434073	Homo sapiens	cDNA FLJ12531 fis, clone NT2RM4000199.	1871	99
119	AAM00936	Homo sapiens	Human bone marrow protein, SEQ ID NO: 412.	3350	100
119	AAB42828	Homo sapiens	Human ORFX ORF2592 polypeptide sequence SEQ ID NO:5184.	2064	100
119	gi9557949	Homo sapiens	mRNA for hypothetical protein (ORF1), clone Telethon(Italy_B41)_Strait02270_FL1 42.	1931	100
120	AAB11082	Homo sapiens	Human secreted protein ZALPHA13 protein.	2783	93
120	gi11230043	Homo sapiens	unnamed protein product	2783	93
120	AAB37988	Homo sapiens	Human secreted protein encoded by gene 5 clone HDPAS92.	2747	93
121	gi12852526	Mus musculus	putative	1689	80
121	AAB41765	Homo sapiens	Human ORFX ORF1529 polypeptide sequence SEQ ID NO:3058.	1576	100
121	gi4406663	Homo sapiens	clone 24945 mRNA sequence, partial cds.	1576	100
122	AAR22958	Homo sapiens	Human proteasome component HC5.	1010	85
122	gi220026	Homo sapiens	Human mRNA for proteasome subunit HC5.	1010	85
122	gi3790135	Homo sapiens	Human DNA sequence from clone RP1-191N21 on chromosome 6q27. Contains a 7 transmembrane receptor (rhodopsin family) (olfactory receptor like) pseudogene, the PDCD2 gene for programmed cell death 2 (RP8 homolog), the TBP gene for TATA box binding protein, the gene for proteasome subunit HC5, ESTs, STSs and GSSs, complete sequence.	1010	85
123	AAB21027	Homo sapiens	Human nucleic acid-binding protein, NuABP-31.	1456	100
123	AAB45146	Homo sapiens	Human secreted protein sequence encoded by gene 27 SEQ ID NO:87.	1456	100
123	gi4884258	Homo sapiens	mRNA; cDNA DKFZp564O092 (from clone DKFZp564O092); partial cds.	1430	100
124	gi13325436	Homo sapiens	Similar to RIKEN cDNA	1394	100

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
			C330013D18 gene, clone MGC:11226 IMAGE:3937599, mRNA, complete cds.		
124	gi13559363	Homo sapiens	MRPL9 mRNA for mitochondrial ribosomal protein L9 (L9mt), complete cds.	1388	99
124	AAG93251	Homo sapiens	Human protein HP02612.	1153	86
125	AAB85507	Homo sapiens	Human protein kinase SGK164.	2949	100
125	gi13543922	Homo sapiens	Similar to RIKEN cDNA 5430416A05 gene, clone MGC:12903 IMAGE:3537086, mRNA, complete cds.	2913	100
125	gi12856491	Mus musculus	putative	2135	.79
126	gi12653817	Homo sapiens	Similar to Male-specific RNA 84Dd, clone MGC:3092 IMAGE:3349383, mRNA, complete cds.	3399	100
126	AAB94115	Homo sapiens	Human protein sequence SEQ ID NO:14356.	3392	99
126	gi10434102	Homo sapiens	cDNA FLJ12549 fis, clone NT2RM4000689.	3392	99
127	gi7243187	Homo sapiens	mRNA for KIAA1403 protein, partial cds.	6448	98
127	gi12652971	Homo sapiens	clone MGC:858 IMAGE:3357380, mRNA, complete cds.	3992	100
127	AAB92872	Homo sapiens	Human protein sequence SEQ ID NO:11460.	3987	99
128	AAB94324	Homo sapiens	Human protein sequence SEQ ID NO:14807.	1779	99
128	gi10434528	Homo sapiens	cDNA FLJ12816 fis, clone NT2RP2002609, weakly similar to 2- HYDROXYMUCONIC SEMIALDEHYDE HYDROLASE (EC 3.1.1).	1779	99
128	AAB42143	Homo sapiens	Human ORFX ORF1907 polypeptide sequence SEQ ID NO:3814.	1521	100
129	gi6329945	Homo sapiens	mRNA for KIAA1140 protein, partial cds.	1857	52
129	gi12805043	Homo sapiens	clone IMAGE:3461487, mRNA, partial cds.	1279	54
129	gi7302173	Drosophila melanogaster	BcDNA:LD21719 gene product	1261	35
130	AAB28199	Homo sapiens	Human HMG-17 non histone chromosomal protein.	322	75
130	gi306864	Homo sapiens	Human non-histone chromosomal protein HMG-17 mRNA, complete cds.	322	75
130	gi32329	Homo sapiens	Human HMG-17 gene for non-histone chromosomal protein HMG-17.	322	75
131	gi16041794	Homo sapiens	clone MGC:23591 IMAGE:4856946, mRNA, complete cds.	2714	99
131	gi15559462	Homo sapiens	Similar to old astrocyte specifically induced substance, clone MGC:20215 IMAGE:4546950, mRNA, complete cds.	2709	99

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
131	gi4519621	Mus musculus	OASIS protein	2406	91
132	gi7573591	Homo sapiens	Human DNA sequence from clone RP1-309K20 on chromosome 20 Contains the gene for a novel protein similar to dysferlin, the SPAG4 gene for sperm associated antigen 4, the CPNE1 gene for Copine I (similar to KIAA0636), the gene KIAA0765 (HRIHFB2091) for an RNA recognition motif (RNP, RRM or RBD domain) containing protein and the 3'	4972	100
132	gi15559252	Homo sapiens	end of the NIFS gene for cysteine desulfurase. Contains ESTs, STSs, GSSs and four putative CpG islands, complete sequence. RNA binding motif protein 12, clone	4972	100
	,	_	MGC:19528 IMAGE:3845090, mRNA, complete cds.		
132	gi15215375	Homo sapiens	RNA binding motif protein 12, clone MGC:16487 IMAGE:3956772, mRNA, complete cds.	4972	100
133	gi12697774	Mus musculus	acetyl-CoA synthetase 2	3181	87
133	gi12697772	Bos taurus	acetyl-CoA synthetase 2	3056	83
133	AAB34712	Homo sapiens	Human secreted protein encoded by DNA clone vo9 1.	2721	100
134	gi7020783	Homo sapiens	cDNA FLJ20580 fis, clone REC00516.	848	100
134	gi15012026	Homo sapiens	Similar to hypothetical protein FLJ20580, clone MGC:13430 IMAGE:4093763, mRNA, complete cds.	848	100
134	gi12833008	Mus musculus	putative	814	85
135	AAB94473	Homo sapiens	Human protein sequence SEQ ID NO:15139.	1970	100
135	AAG74880	Homo sapiens	Human colon cancer antigen protein SEQ ID NO:5644.	1970	100
135	AAB43720	Homo sapiens	Human cancer associated protein sequence SEQ ID NO:1165.	1970	100
136	gi10047285	Homo sapiens	mRNA for KIAA1605 protein, partial cds.	3610	99
136	gi16215453	Homo sapiens	mRNA for bile acid beta-glucosidase.	3610	99
136	gi15030210	Homo sapiens	KIAA1605 protein, clone MGC:16895 IMAGE:4339156, mRNA, complete cds.	3610	99
137	gi4914601	Homo sapiens	mRNA; cDNA DKFZp564A026 (from clone DKFZp564A026).	4171	94
137	AAB94357	Homo sapiens	Human protein sequence SEQ ID NO:14881.	2195	99
137	AAY45161	Homo sapiens	Human secreted protein clone CO139_3 protein sequence.	2112	100
138	gi313131	Torpedo marmorata	alpha-tubulin	1192	97
138	gi14198110	Mus musculus	tubulin alpha 1	1192	97
138	gi13435777	Mus musculus	tubulin alpha 6	1192	97

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
139	AAB94856	Homo sapiens	Human protein sequence SEQ ID NO:16044.	2138	100
139	AAB94628	Homo sapiens	Human protein sequence SEQ ID NO:15490.	2138	100
139	gi10436294	Homo sapiens	cDNA FLJ13970 fis, clone Y79AA1001533, moderately similar to Mouse mRNA for RNA polymerase I associated factor (PAF53).	2138	100
140	gi2642187	Rattus norvegicus	endo-alpha-D-mannosidase	1415	67
140	AAB95204	Homo sapiens	Human protein sequence SEQ ID NO:17303.	1094	66
140	gi10434559	Homo sapiens	cDNA FLJ12838 fis, clone NT2RP2003230, moderately similar to Rattus norvegicus endo-alpha-D-	1094	66
141	gi3449308	Homo sapiens	mannosidase (Enman) mRNA. mRNA for MEGF8, partial cds.	9785	100
141	gi6681364	Rattus norvegicus	MEGF8	4772	95
141	gi10728654	Drosophila melanogaster	CG7466 gene product	2902	34
142	AAY29517	Homo sapiens	Human lung tumour protein SAL-82 predicted amino acid sequence.	3048	100
142	gi13958036	Homo sapiens	FYVE-finger protein EIP1 mRNA, complete cds.	3048	100
142	AAY29861	Homo sapiens	Human secreted protein clone cb98 4.	3041	99
143	gi14718539	Homo sapiens	HIC-3 mRNA, complete cds.	3178	99
143	gi5689371	Homo sapiens	mRNA for KIAA1020 protein, partial cds.	2970	99
143	gi7328028	Homo sapiens	mRNA; cDNA DKFZp434F0616 (from clone DKFZp434F0616); partial cds.	1738	100
144	gi12620400	Homo sapiens	mitochondrial carrier protein CGI-69 long form mRNA, complete cds.	1856	99
144	AAB42783	Homo sapiens	Human ORFX ORF2547 polypeptide sequence SEQ ID NO:5094.	1804	96
144	gi10438783	Homo sapiens	cDNA: FLJ22407 fis, clone HRC08407.	1798	97
145	gi2792366	Homo sapiens	unknown protein IT12 mRNA, partial cds.	4390	99
145	gi1843399	Homo sapiens	mRNA, partial cds, clone:RES4-25.	3676	99
145	gi14602505	Homo sapiens	clone IMAGE:3936655, mRNA, partial cds.	2366	99
146	gi13359167	Homo sapiens	mRNA for KIAA1646 protein, partial cds.	2581	99
146	AAY96059	Homo sapiens	Human sphingosine kinase C.	2456	99
146	gi6572330	Homo sapiens	Human DNA sequence from clone 59H18 on chromosome 22. Contains the 3' part of the gene for KIAA0767, a novel gene, ESTs, STSs, GSSs and a putative CpG island, complete sequence.	1627	96
147	gi14043303	Homo sapiens	exonuclease NEF-sp, clone MGC:15944 IMAGE:3537866, mRNA,	4043	100

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
			complete cds.	 	
147	gi13272524	Homo sapiens	exonuclease NEF-sp mRNA, complete cds.	4039	99
147	gi12053043	Homo sapiens	mRNA; cDNA DKFZp434J0315 (from clone DKFZp434J0315); complete cds.	3843	95
148	gi7243037	Homo sapiens	mRNA for KIAA1328 protein, partial cds.	2894	100
148	gi13874541	Macaca fascicularis	hypothetical protein	2492	93
148	gi1335313	Homo sapiens	Human muscle mRNA for embryonic myosin heavy chain (SMHCE).	129	24
149	AAB42399	Homo sapiens	Human ORFX ORF2163 polypeptide sequence SEQ ID NO:4326.	1362	91
149	AAB42366	Homo sapiens	Human ORFX ORF2130 polypeptide sequence SEQ ID NO:4260.	626	100
149	gi7298594	Drosophila melanogaster	CG10189 gene product	223	35
150	AAB95372	Homo sapiens	Human protein sequence SEQ ID NO:17692.	1538	99
150	gi10435150	Homo sapiens	cDNA FLJ13220 fis, clone NT2RP4002047, moderately similar to GTP-BINDING PROTEIN LEPA.	1538	99
150	gi10437720	Homo sapiens	cDNA: FLJ21595 fis, clone COL07069.	1438	100
151	gi3327080	Homo sapiens	mRNA for KIAA0633 protein, partial cds.	6823	99
151	gi857571	Mus musculus	cordon-bleu gene product	1345	81
151	gi6094680	Homo sapiens	PAC clone RP5-1168M19 from 7p12- q11.21, complete sequence.	1342	100
152	gi15451265	Macaca fascicularis	hypothetical protein	2728	98
152	AAB41597	Homo sapiens	Human ORFX ORF1361 polypeptide sequence SEQ ID NO:2722.	2650	100
152	gi5689443	Homo sapiens	mRNA for KIAA1053 protein, partial cds.	2650	100
153	gi14036062	Homo sapiens	unnamed protein product	1930	100
153	AAG81377	Homo sapiens	Human AFP protein sequence SEQ ID NO:272.	1925	99
153	gi12833112	Mus musculus	putative	1727	88
154	gi12832455	Mus musculus	putative	1220	89
154	gi15080314	Homo sapiens	Similar to RIKEN cDNA 0610010D20 gene, clone MGC:20590 IMAGE:4310241, mRNA, complete cds.	514	100
154	gi6002488	Penicillium chrysogenum	hypothetical protein	338	31
155	gi14017889	Homo sapiens	mRNA for KIAA1836 protein, partial cds.	2511	100
155	AAB94592	Homo sapiens	Human protein sequence SEQ ID NO:15402.	972	50
155	gi10435321	Homo sapiens	cDNA FLJ13337 fis, clone OVARC1001880.	972	50
156	gi14550510	Homo sapiens	pseudouridylate synthase 1, clone	2123	100

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
			MGC:2736 IMAGE:2822709, mRNA, complete cds.		
156	gi12804097	Homo sapiens	Similar to pseudouridine synthase 1, clone MGC:11268 IMAGE:3943243, mRNA, complete cds.	2123	100
156	gi4455035	Homo sapiens	pseudouridine synthase 1 (PUS1) mRNA, partial cds.	1927	99
157	AAY58052	Homo sapiens	Human protein kinase H2LAU20 protein sequence.	3198	98
157	gi9652080	Homo sapiens	protein kinase DYRK4 (DYRK4) mRNA, partial cds.	2844	100
157	AAW71685	Homo sapiens	Amino acid sequence of human serine/threonine protein kinase.	1909	97
158	gi7300952	Drosophila melanogaster	BcDNA:LD21504 gene product	971	62
158	gi4972728	Drosophila melanogaster	unknown	971	62
158	AAB97646	Homo sapiens	Ribosomal S3 protein 17.	831	99
159	AAU02201	Homo sapiens	Phosphatase 1 protein-like protein, MEM6.	1514	100
159	gi15551577	Homo sapiens	unnamed protein product	1514	100
159	AAB95633	Homo sapiens	Human protein sequence SEQ ID NO:18363.	1510	99
160	gi12804573	Homo sapiens	Similar to CG11334 gene product, clone MGC:3207 IMAGE:3501899, mRNA, complete cds.	1859	100
160	gi12851419	Mus musculus	putative	1590	86
160	gi7302053	Drosophila melanogaster	CG11334 gene product	1046	59
161	gi1580781	Homo sapiens	Human beige-like protein (BGL) mRNA, partial cds.	9734	99
161	gi10180266	Mus musculus	LBA	9333	86
161	gi10257401	Mus musculus	LBA isoform beta	8920	86
162	gi15082589	Homo sapiens	clone MGC:4408 IMAGE:2906200, mRNA, complete cds.	2065	99
162	gi15638615	Arabidopsis thaliana	HEN1	350	37
162	gi13241746	Arabidopsis thaliana	CORYMBOSA2	350	37
163	gi15291227	Drosophila melanogaster	GH13040p	701	40
163	gi7303780	Drosophila melanogaster	CG12214 gene product	701	40
163	AAB95882	Homo sapiens	Human protein sequence SEQ ID NO:18991.	501	100
164	gi3327170	Homo sapiens	mRNA for KIAA0678 protein, partial cds.	5255	100
164	AAB95304 ·	Homo sapiens	Human protein sequence SEQ ID NO:17542.	4431	99
164	gi14134120	Caenorhabditis elegans	endocytosis protein RME-8	2127	42
165	AAB53427	Homo sapiens	Human colon cancer antigen protein sequence SEQ ID NO:967.	813	96

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
165	gi13905098	Mus musculus	B-cell translocation gene 1, anti- proliferative	813	96
165	gi293306	Mus musculus	B-cell translocation gene-1 protein	813	96
166	gi13365897	Macaca fascicularis	hypothetical protein	2501	97
166	AAY02168	Homo sapiens	A facilitative glucose transporter protein GLUT8.	870	99
166	gi13445575	Homo sapiens	facilitative glucose transporter GLUT10 (SLC2A10) mRNA, complete cds.	835	39
167	gi13365897	Macaca fascicularis	hypothetical protein	2173	97
167	AAY02168	Homo sapiens	A facilitative glucose transporter protein GLUT8.	870	99
167	gi13445575	Homo sapiens	facilitative glucose transporter GLUT10 (SLC2A10) mRNA, complete cds.	678	37
168	gi10047251	Homo sapiens	mRNA for KIAA1588 protein, partial cds.	3292	100
168	gi14424704	Homo sapiens	clone MGC:15071 IMAGE:4110510, mRNA, complete cds.	2315	100
168	gi4567179	Homo sapiens	chromosome 19, BAC 37295 (CIT-B-21A4), complete sequence.	1269	43
169	gi15558943	Homo sapiens	guanylate binding protein 4 mRNA, complete cds.	3134	99
169	gi1174187	Mus musculus	purine nucleotide binding protein	2260	70
169	gi193444	Mus musculus	guanylate binding protein	1986	66
170	gi14585859	Homo sapiens	hypothetical protein SB138	1121	100
170	gi6665778	Mus musculus	cyclin ania-6b	1052	92
170	gi12841169	Mus musculus	putative	1052	92
171	AAB64407	Homo sapiens	Amino acid sequence of human intracellular signalling molecule INTRA39.	3394	100
171	AAB71963	Homo sapiens	Human TGF-beta receptor encoded by cDNA clone HFIHY04.	3394	100
171	gi10438113	Homo sapiens	cDNA: FLJ21908 fis, clone HEP03830.	3385	99
172	gi12652533	Homo sapiens	clone MGC:2637 IMAGE:3505128, mRNA, complete cds.	676	89
172	AAB67453	Homo sapiens	Amino acid sequence of a human chaperone polypeptide.	668	88
172	gi9758421	Arabidopsis thaliana	gene_id:MHF15.7~similar to unknown protein~	199	28
173	AAB97025	Homo sapiens	Human colon carcinoma suppressor gene-related protein.	1773	61
173	gi9857318	Homo sapiens	Asef mRNA for APC-stimulated guanine nucleotide exchange factor, complete cds.	1773	61
173	gi8809845	Homo sapiens	chromosome 2q22 RhoGEF mRNA, complete cds.	1700	61
174	gi12052828	Homo sapiens	mRNA; cDNA DKFZp564N1062 (from clone DKFZp564N1062); complete cds.	1601	99
174	gi12850603	Mus musculus	putative	1062	92

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
174	AAB94655	Homo sapiens	Human protein sequence SEQ ID NO:15568.	671	100
175	gi15080282	Homo sapiens	Similar to putative sialoglycoprotease type 2, clone MGC:20293 IMAGE:4121450, mRNA, complete cds.	1747	99
175	gi11071727	Homo sapiens	mRNA for putative sialoglycoprotease type 2.	1707	92
175	gi12847276	Mus musculus	putative	1541	84
176	AAB36628	Homo sapiens	Human FLEXHT-50 protein sequence SEQ ID NO:50.	527	100
176	AAB94208	Homo sapiens	Human protein sequence SEQ ID NO:14557.	527.	100
176	AAG01512	Homo sapiens	Human secreted protein, SEQ ID NO: 5593.	527	100
177	gi15929052	Homo sapiens	Similar to RIKEN cDNA 2810442O16 gene, clone MGC:23197 IMAGE:4861869, mRNA, complete cds.	2084	100
177	gi11493155	Homo sapiens	Human DNA sequence from clone RP5-852M4 on chromosome 20. Contains the gene encoding the HBV associated factor, a novel gene similar to Drosophilia CG17883, a putative novel gene, two CpG islands, ESTs, GSSs, and STSs, complete sequence.	1952	100
177	gi12840168	Mus musculus	putative	1938	93
178	AAB87034	Homo sapiens	Human secreted protein TANGO 339, SEQ ID NO:3.	1449	100
178	AAY76266	Homo sapiens	Human secreted protein encoded by gene 10 fragment.	1449	100
178	AAB87135	Homo sapiens	Human secreted protein TANGO 339 F20Y variant, SEQ ID NO:139.	1446	99
179	gi434763	Homo sapiens	Human mRNA for KIAA0120 gene, complete cds.	1048	100
179	gi14424677	Homo sapiens	transgelin 2, clone MGC:15279 IMAGE:4301018, mRNA, complete cds.	1048	100
179	gi9956026	Homo sapiens	clone CDABP0035 mRNA sequence.	1048	100
180	AAB31677	Homo sapiens	Amino acid sequence of a human protein having a hydrophobic domain.	2803	100
180	AAE03346	Homo sapiens	Human gene 19 encoded secreted protein HCRNF14, SEQ ID NO:120.	2803	100
180	AAE03310	Homo sapiens	Human gene 19 encoded secreted protein HCRNF14, SEQ ID NO:84.	2803	100
181	AAB41910	Homo sapiens	Human ORFX ORF1674 polypeptide sequence SEQ ID NO:3348.	1530	99
181	gi5262467	Homo sapiens	mRNA; cDNA DKFZp564I122 (from clone DKFZp564I122).	1530	99
181	gi12849716	Mus musculus	putative	1259	82
182	gi2072972	Homo sapiens	Human L1 element L1.25 p40 and putative p150 genes, complete cds.	497	53
182	AAB64943	Homo sapiens	Human secreted protein sequence	494	54

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
			encoded by gene 7 SEQ ID NO:121.		
182	gi5070622	Homo sapiens	retrotransposon L1 insertion in X- linked retinitis pigmentosa locus, complete sequence.	494	53
183	AAB59191	Homo sapiens	Human NADE.	217	47
183	gi8452894	Homo sapiens	p75NTR-associated cell death executor (NADE) mRNA, complete cds.	217	47
183	gi189379	Homo sapiens	Human unknown protein from clone pHGR74 mRNA, complete cds.	217	47
184	AAB88468	Homo sapiens	Human membrane or secretory protein clone PSEC0263.	4931	97
184	gi14272788	Homo sapiens	unnamed protein product	4931	97
184	gi577301	Homo sapiens	Human mRNA for KIAA0090 gene, partial cds.	4650	99
185	AAG64953	Homo sapiens	Human ATP-dependent helicase protein 68.	3169	100
185	gi12052748	Homo sapiens	mRNA; cDNA DKFZp564B1023 (from clone DKFZp564B1023); complete cds.	2716	100
185	gi12836314	Mus musculus	putative	2655	83
186	gi14017781	Homo sapiens	mRNA for KIAA1782 protein, partial cds.	2834	99
186	gi4062983	Mus musculus	Eos protein	2747	95
186	gi11612390	Homo sapiens	zinc finger transcription factor Eos mRNA, complete cds.	2603	98
187	AAB95721	Homo sapiens	Human protein sequence SEQ ID NO:18592.	2419	100
187	gi10436538	Homo sapiens	cDNA FLJ14153 fis, clone NT2RM1000092, weakly similar to MULTIDRUG RESISTANCE PROTEIN 2.	2419	100
187	gi12248763	Homo sapiens	mRNA for SMAP-4, complete cds.	2323	96
188	gi13278906	Homo sapiens	clone MGC:4440 IMAGE:2959536, mRNA, complete cds.	1040	100
188	gi13278819	Homo sapiens	clone MGC:2776 IMAGE:2959536, mRNA, complete cds.	1040	100
188 .	AAB95829	Homo sapiens	Human protein sequence SEQ ID NO:18847.	618	79
189	gi14602977	Homo sapiens	Similar to KIAA0789 gene product, clone MGC:16602 IMAGE:4110708, mRNA, complete cds.	3100	99
189	gi3043570	Homo sapiens	mRNA for KIAA0523 protein, partial cds.	2564	100
189	gi14133217	Homo sapiens	mRNA for KIAA0789 protein, partial cds.	1463	49
190	gi9717245	Mus musculus	cytoplasmic dynein heavy chain	5569	98
190	gi294543	Rattus norvegicus	dynein heavy chain	5557	98
190	gi402528	Rattus norvegicus	cytoplasmic dynein heavy chain	5535	98
191	gi13537204	Homo sapiens	mRNA for MAST205, complete cds.	6834	98
191	gi406058	Mus musculus	protein kinase	6343	86
191	gi3882335	Homo sapiens	mRNA for KIAA0807 protein, partial	6300	98

Table 2

		<u></u>	1	1	Identity
			cds.	 	1
192	gi12847109	Mus musculus	putative	1356	79
192	gi13623271	Homo sapiens	Similar to RIKEN cDNA 2600005P05 gene, clone MGC:11321 IMAGE:3951804, mRNA, complete cds.	1332	100
192	gi12847837	Mus musculus	putative	1170	76
193	gi38149	Pongo pygmaeus	epsilon-globin	397	100
193	gi903731	Gorilla gorilla	epsilon-globin	397	100
193	gi903707	Pan troglodytes	epsilon-globin	397	100
194	AAB74695	Homo sapiens	Human membrane associated protein MEMAP-1.	1799	100
194	AAE01340	Homo sapiens	Human gene 22 encoded secreted protein fragment, SEQ ID NO:205.	1799	100
194	gi15929183	Homo sapiens	modulator of apoptosis 1, clone MGC:9487 IMAGE:3922055, mRNA, complete cds.	1799	100
195	AAG93260	Homo sapiens	Human protein HP10106.	1769	100
195	gi15029765	Mus musculus	RIKEN cDNA 2810039M17 gene	1650	91
195	gi12849932	Mus musculus	putative	1650	91
196	gi14017843	Homo sapiens	mRNA for KIAA1813 protein, partial cds.	3434	100
196	gi15193290	Homo sapiens	LAPSER1 (LAPSER1) mRNA, complete cds.	3309	100
196	gi8217421	Homo sapiens	Human DNA sequence from clone RP11-108L7 on chromosome 10. contains part of the gene for a novel Insulin-like growth factor binding type protein with Kazal-type serine protease inhibitor domain, the gene for a novel protein similar to rat tricarboxylate carrier, the gene for a novel PDZ (DHR, GLGF) domain protein, the gene for a novel protein similar to KIAA0552, KIAA0341 and Fugu hypothetical protein 2, the gene for a novel protein similar to Plasmodium POM1 and C. elegans F46G11.1, a putative novel gene, the SEMA4G gene for semaphorin 4G and a novel gene. Contains ESTs, STSs, GSSs and seven putative CpG islands, complete sequence.	3264	100
197	gi1458241	Caenorhabditis elegans	Hypothetical protein B0507.2	782 .	39
197	gi12832510	Mus musculus	putative	490	89
197	AAB54014	Homo sapiens	Human pancreatic cancer antigen protein sequence SEQ ID NO:466.	242	100
198	gi500747	Mus musculus	capping protein beta-subunit, isoform 1	1440	98
198	gi212902	Galius gallus	actin-capping protein Z beta subunit		

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
			Z-line, beta		
199	gi14017787	Homo sapiens	mRNA for KIAA1785 protein, partial cds.	3195	100
199	gi13436428	Homo sapiens	Similar to feminization 1 a homolog (C. elegans), clone MGC:4216 IMAGE:2957950, mRNA, complete cds.	2197	64
199	gi12836689	Mus musculus	putative	2164	65
200	gi7959811	Homo sapiens	PRO1167	389	100
200	gi2736345	Caenorhabditis elegans	contains similarity to G-coupled protein receptors	69	33
200	gi7504953	Caenorhabditis elegans	hypothetical protein H22D07.1 - Caenorhabditis elegans >	69	33
201	gi12697975	Homo sapiens	mRNA for KIAA1715 protein, partial cds.	2230	100
201	AAB42461	Homo sapiens	Human ORFX ORF2225 polypeptide sequence SEQ ID NO:4450.	1015	100
201	gi12844031	Mus musculus	putative	567	92
202	gi7296176	Drosophila melanogaster	CG2839 gene product	195	27
202	gi10438900	Homo sapiens	cDNA: FLJ22490 fis, clone HRC10983.	184	97
202	gi5824430	Caenorhabditis elegans	cDNA EST yk501h2.5 comes from this gene~cDNA EST yk523d4.5 comes from this gene~cDNA EST yk553f6.5 comes from this gene~cDNA EST yk595g12.5 comes from this gene~cDNA EST yk606g10.5 comes from this gene~cDNA EST yk652f3.5 comes from this gene	182	21
203	AAM00957	Homo sapiens	Human bone marrow protein, SEQ ID NO: 433.	1725	100
203	gi4151807	Rattus norvegicus	membrane-associated guanylate kinase- interacting protein 2 Maguin-2	1484	62
203	gi4151805	Rattus norvegicus	membrane-associated guanylate kinase- interacting protein 1 Maguin-1	1484	62
204	AAM00844	Homo sapiens	Human bone marrow protein, SEQ ID NO: 207.	1051	98
204	gi4151807	Rattus norvegicus	membrane-associated guanylate kinase- interacting protein 2 Maguin-2	779	69
204	gi4151805	Rattus norvegicus	membrane-associated guanylate kinase- interacting protein 1 Maguin-1	779	69
205	AAM00957	Homo sapiens	Human bone marrow protein, SEQ ID NO: 433.	1576	92
205	gi4151807	Rattus norvegicus	membrane-associated guanylate kinase- interacting protein 2 Maguin-2	1349	57
205	gi4151805	Rattus norvegicus	membrane-associated guanylate kinase- interacting protein 1 Maguin-1	1349	57 ·
206	gi7242969	Homo sapiens	mRNA for KIAA1307 protein, partial cds.	8582	99
206	AAM00860	Homo sapiens	Human bone marrow protein, SEQ ID NO: 223.	4841	98
206	gi4426611	Drosophila	pushover	2137	46

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
		melanogaster			1 200,000
207	AAB62210	Homo sapiens	Human ABCA2 transporter protein.	9835	99
207	gi13173186	Homo sapiens	ABC transporter ABCA2 (ABCA2) mRNA, complete cds.	9835	99
207	gi9957467	Homo sapiens	ATP-binding cassette sub-family A member 2 (ABCA2) mRNA, complete cds.	9835	99
208	AAB94358	Homo sapiens	Human protein sequence SEQ ID NO:14883.	2268	99
208	gi10434632	Homo sapiens	cDNA FLJ12886 fis, clone NT2RP2004041, weakly similar to SYNAPSINS IA AND IB.	2268	99
208	gi12052738	Homo sapiens	mRNA; cDNA DKFZp564H1322 (from clone DKFZp564H1322); complete cds.	2268	99
209	gi14627122	Homo sapiens	Human DNA sequence from clone RP4-583P15 on chromosome 20 Contains ESTs, STSs, GSSs and ten CpG islands. Contains the TNFRSF6B gene for tumor necrosis factor receptor 6b (decoy), the 3' part of the	2074	99
	-		KIAA1088 gene, the ARFRP1 gene for ADP-ribosylation factor related protein 1, two genes for novel proteins, the		
• .			gene for a GLUT4 enhancer factor and the gene for a novel zinc finger protein similar to rat RIN ZF and the gene for a novel BTB/POZ domain containing		
209	gi13162677	Homo sapiens	zinc finger protein, complete sequence. GLUT4 enhancer factor mRNA, complete cds.	2055	98
209	gi12655101	Homo sapiens	clone IMAGE:3140406, mRNA, partial cds.	1766	100
210	gi14279329	Homo sapiens	ubiquitin specific protease (USP28) mRNA, complete cds.	4131	92
210	gi7959297	Homo sapiens	mRNA for KIAA1515 protein, partial cds.	3872	100
210	AAB31552	Homo sapiens	A human ubiquitin specific protease 25 (USP25).	2058	48
211	AAB36579	Homo sapiens	Human FLEXHT-1 protein sequence SEQ ID NO:1.	1829	100 .
211	AAB94048	Homo sapiens	Human protein sequence SEQ ID NO:14211.	1825	99
211	gi10433984	Homo sapiens	cDNA FLJ12475 fis, clone NT2RM1000962.	1825	99
212	gi15824499	Homo sapiens	GalNAc-4-O-sulfotransferase 1 mRNA, complete cds.	2238	100
212	gi11990885	Homo sapiens	GalNAc4ST mRNA for GalNAc 4- sulfotransferase, complete cds.	2238	100
212	gi15559803	Homo sapiens	carbohydrate (N-acetylgalactosamine 4-0) sulfotransferase 8, clone MGC:20987 IMAGE:4635405, mRNA, complete cds.	2238	100

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
213	AAB43387	Homo sapiens	Human ORFX ORF3151 polypeptide sequence SEQ ID NO:6302.	1056	100
213	gi15292317	Drosophila melanogaster	LD46863p	549	50
213	gi7302029	Drosophila melanogaster	CG12054 gene product	549	50
214	gi12843216	Mus musculus	putative	913	84
214	gi14585867	Homo sapiens	hypothetical protein SB145	297	44
214	gi14388386	Macaca fascicularis	hypothetical protein	295	44
215	gi14133219	Homo sapiens	mRNA for KIAA0833 protein, partial cds.	7195	99
215	gi6580410	Homo sapiens	Human DNA sequence from clone RP3-467L1 on chromosome 1p36.21-36.33. Contains the 3' part of gene KIAA0833, the VAMP3 gene for vesicle-associated membrane protein 3 (cellubrevin), the PER3 gene for period (Drosophila) homolog 3 and the gene for urotensin II. Contains two putative CpG islands, ESTs, STSs and GSSs, complete sequence.	3642	99
215	AAB42729	Homo sapiens	Human ORFX ORF2493 polypeptide sequence SEQ ID NO:4986.	997	54
216	gi7293088	Drosophila melanogaster	CG9213 gene product	811	30
216	gi15810333	Arabidopsis thaliana	unknown protein	713	28
216	gi13324888	Caenorhabditis elegans	Hypothetical protein B0361.2	710	34
217	gi2443331	Xenopus laevis	Nfrl .	2421	75
217	AAB34944	Homo sapiens	Human secreted protein sequence encoded by gene 20 SEQ ID NO:148.	1129	91
217	gi15292543	Drosophila melanogaster	SD06560p	911	36
218	gi7243111	Homo sapiens	mRNA for KIAA1365 protein, partial cds.	3855	100
218	gi1657758	Rattus norvegicus	densin-180	3640	93
218	gi8570180	Rattus norvegicus	densin-180 variant D	1250	83
219	gi14017839	Homo sapiens	mRNA for KIAA1811 protein, partial cds.	1726	80
219	gi3217028	Homo sapiens	mRNA for putative serine/threonine protein kinase, partial.	1450	84
219	gi7294217	Drosophila melanogaster	CG6114 gene product	1055	70
220	gi7297674	Drosophila melanogaster	CG13139 gene product	942	75
220	gi12857050	Mus musculus	putative	767	62
220	gi15636900	Gallus gallus	avEna neural variant	139	52
221	gi15489242	Homo sapiens	clone IMAGE:3859726, mRNA,	1001	88

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
			partial cds.	 	1
221	gi13543991	Homo sapiens	clone IMAGE:3627860, mRNA, partial cds.	1001	88
221	gi12847182	Mus musculus	putative	328	39
222	gi14133209	Homo sapiens	mRNA for KIAA0654 protein, partial cds.	6089	99
222	gi930343	Homo sapiens	Human LAR-interacting protein 1b mRNA, complete cds.	3559	60
222	gi930341	Homo sapiens	Human LAR-interacting protein 1a mRNA, complete cds.	3503	60
223	gi12620207	Homo sapiens	Clorf25 mRNA, complete cds.	3807	98
223	gi9588430	Homo sapiens	Human DNA sequence from clone GS1-120K12 on chromosome 1q25.3-31.2. Contains the gene for ring finger protein DING or BAP-1, an FTH1 (ferritin, heavy polypeptide 1) pseudogene, the 3' end of the gene for a novel protein similar to archaeal, yeast and worm N2,N2-dimethylguanosine tRNA methyltransferase, ESTs, STSs, GSSs and two putative CpG islands, complete sequence.	2300	98
223	gi12835704	Mus musculus	putative	1420	00
224	gi14595658	Xenopus laevis	LIM protein prickle	2865	67
224	gi10727796	Drosophila melanogaster	esn gene product	698	42
224	gi6634092	Drosophila melanogaster	LIM-domain protein	698	42
225	gi13375149	Homo sapiens	Human DNA sequence from clone RP5-1118M15 on chromosome 20 Contains part of a gene similar to P14 Bos taurus (P14L), a novel gene, ESTs, STSs, GSSs and a CpG Island, complete sequence.	957	99
225	gi7259265	Mus musculus	contains transmembrane (TM) region	314	
225	AAY53871	Homo sapiens	A human brain-derived signalling factor polypeptide.	299	50 45
226	gi12803987	Homo sapiens	clone MGC:4174 IMAGE:3634226, mRNA, complete cds.	743	100
226	gi12805417	Mus musculus	Unknown (protein for MGC:7354)	444	66
226	gi12849498	Mus musculus	putative	235	72
227	AAY91629	Homo sapiens	Human secreted protein sequence encoded by gene 23 SEQ ID NO:302.	1391	87
227	gi7677403	Homo sapiens	F-box protein FBG2 (FBG2) mRNA, complete cds.	1391	87
227	AAY83046	Homo sapiens	F-box protein FBP-6.	1333	82
228	gi15079958	Homo sapiens	chromosome 11 open reading frame 24, clone MGC:19741 IMAGE:3614861, mRNA, complete cds.	2231	99
228	gi11527205	Homo sapiens	DM4E3 (Cl lorf24) mRNA, complete cds.	2224	99

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Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
228	AAB18965	Homo sapiens	Amino acid sequence of a human transmembrane protein.	2055	99
229	gi15930199	Homo sapiens	Similar to RIKEN cDNA 4921523I18 gene, clone MGC:9467 IMAGE:3914747, mRNA, complete cds.	1451	99
229	gi13278594	Mus musculus	RIKEN cDNA 4921523I18 gene	1440	97
229	gi12856904	Mus musculus	putative	1440	97
230	gi15680131	Homo sapiens	hypothetical protein FLJ12171, clone MGC:19889 IMAGE:4652087, mRNA, complete cds.	1638	100
230	gi14043242	Homo sapiens	hypothetical protein FLJ12171, clone MGC:15694 IMAGE:3351601, mRNA, complete cds.	1638	100
230	AAB93912	Homo sapiens	Human protein sequence SEQ ID NO:13880.	1634	99
231	AAB56947	Homo sapiens	Human prostate cancer antigen protein sequence SEQ ID NO:1525.	779	100
231	AAB68408	Homo sapiens	Amino acid sequence of a human NOV1 polypeptide.	574	100
231	AAY81695	Homo sapiens	Human PTN protein sequence.	574	100
232	gi11138034	Homo sapiens	mRNA for KIAA1173 protein, complete cds.	2665	100
232	AAG89259	Homo sapiens	Human secreted protein, SEQ ID NO: 379.	2654	99
232	gi12834372	Mus musculus	putative	2427	90
233	AAB98612	Homo sapiens	Human tumour suppressor gene, TSG16, protein.	1706	55
233	gi11596412	Homo sapiens	GAC-1 (GAC-1) mRNA, complete cds.	893	77
233	gi4240237	Homo sapiens	mRNA for KIAA0874 protein, partial cds.	893	77
234	AAB41108	Homo sapiens	Human ORFX ORF872 polypeptide sequence SEQ ID NO:1744.	4170	99
234	gi6331287	Homo sapiens	mRNA for KIAA1274 protein, partial cds.	3936	99
234	gi1545959	Mus musculus	paladin	3560	80
235	gi9368849	Homo sapiens	mRNA; cDNA DKFZp761G2113 (from clone DKFZp761G2113).	972	99
235	gi7293878	Drosophila melanogaster	CG13379 gene product	274	36
235	gi14532482	Arabidopsis thaliana	AT5g58570/mzn1_20	152	31
236	gi3242242	Mus musculus	hyperpolarization-activated cation channel, HAC2	4309	91
236	gi7407645	Rattus norvegicus	hyperpolarization-activated, cyclic nucleotide-gated potassium channel 1	4306	91
236	gi2708316	Mus musculus	brain cyclic nucleotide gated 1; Bcng- 1; brain specific ion channel protein	4301	91
237	AAB13370	Homo sapiens	Human brain-associated protein HBAP-1.	1055	100
237	gi9944291	Homo sapiens	TTYH1 mRNA, complete cds.	1055	100
237	gi9651109	Macaca fascicularis	ТТҮНІ	1032	98

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
238	AAU00476	Homo sapiens	Human INTERCEPT 400 protein.	1428	100
238	AAY79266	Homo sapiens	Human elongase homologue HS3.	1428	100
238	AAB29648	Homo sapiens	Human membrane-associated protein HUMAP-5.	1428	100
239	AAB84885	Homo sapiens	Human protein, SEQ ID 14.	4029	99
239	AAB84882	Homo sapiens	Human protein, SEQ ID 6.	4029	99
239	gi5262593	Homo sapiens	mRNA; cDNA DKFZp434N093 (from clone DKFZp434N093); partial cds.	3684	99
240	gi13477247	Homo sapiens	Similar to RIKEN cDNA 5031400M07 gene, clone MGC:13079 IMAGE:3840918, mRNA, complete cds.	2153	100
240	AAB18987	Homo sapiens	Amino acid sequence of a human transmembrane protein.	2148	.99
240	gi7670425	Mus musculus	unnamed protein product	1904	89
241	AAG63222	Homo sapiens	Amino acid sequence of a human lipid metabolism enzyme.	2194	100
241	gi14861069	Mus musculus	phosphatidyl inositol phosphate kinase type II gamma	2120	95
241	gi3387798	Rattus norvegicus	phosphatidylinositol 5-phosphate 4-kinase gamma	2087	95
242	gi7295732	Drosophila melanogaster	ft gene product	2915	39 ·
242	gi157409	Drosophila melanogaster	fat protein	2901	39
242	gi10727403	Drosophila melanogaster	ds gene product	2236	34
243	AAF90315_aa 2	Homo sapiens	Winged helix/zinc finger transcription factor FOXP1 cDNA.	819	98
243	AAB82339	Homo sapiens	Winged helix/zinc finger transcription factor FOXP1.	819	98
243	gi12043714	Homo sapiens	clone pAB195 FOXP1 (FOXP1) mRNA, complete cds.	819	98
244	gi10440073	Homo sapiens	cDNA: FLJ23399 fis, clone HEP18254.	2620	100
244	gi7018524	Homo sapiens	mRNA; cDNA DKFZp762K137 (from clone DKFZp762K137); partial cds.	2524	100
244	gi14133227	Homo sapiens	mRNA for KIAA0970 protein, partial cds.	1367	51
245	AAB94855	Homo sapiens	Human protein sequence SEQ ID NO:16042.	1347	100
245	gi10436290	Homo sapiens	cDNA FLJ13968 fis, clone Y79AA1001493, weakly similar to UBIQUITIN-CONJUGATING ENZYME E2-17 KD 9 (EC 6.3.2.19).	1347	100
245	gi16198439	Homo sapiens	hypothetical protein FLJ13855, clone MGC:16842 IMAGE:3915698, mRNA, complete cds.	1347	100
246	gi6330302	Homo sapiens	mRNA for KIAA1185 protein, partial cds.	2041	100
246	AAG74603	Homo sapiens	Human colon cancer antigen protein SEQ ID NO:5367.	1530	97
246	AAB53321	Homo sapiens	Human colon cancer antigen protein sequence SEQ ID NO:861.	1530	97

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
247	gi535390	Macronuclear Homo sapiens	Human cellular retinol binding protein II (CRBPII) mRNA, complete cds.	715	99
247	gi397352	Mus musculus	mCRBPII	674	91
247	gi12833902	Mus musculus	putative	669	90
248	AAG01285	Homo sapiens	Human secreted protein, SEQ ID NO: 5366.	209	87
248	AAR05562	Homo sapiens	Laminin -binding protein encoded by insert from J9 lambda gt10 phage.	209	87
248	gi1149509	Gallus gallus	37kD Laminin receptor precursor /p40 ribosomal associated protein	209	87
249	gi13162226	Homo sapiens	Human DNA sequence from clone RP4-543J19 on chromosome 20 Contains part of the GNAS1 gene encoding guanine nucleotide binding protein (G protein, alpha stimulating activity polypeptide 1) including neuroendocrine secretory protein 55 (NESP55), the CTSZA gene encoding cathepsin Z, the ATP5E gene encoding ATP synthase (H+ transporting, mitochondrial F1 complex, epsilon subunit), the gene encoding protein HSPC130 (TH1 Drosophila homolog), the gene for tubulin beta 1 class VI (TUBB1), a gene encoding the CGI-107 protein (LOC51012), four CpG	1591	100
249	gi11230445	Homo sapiens	islands, ESTs, STSs and GSSs, complete sequence. TUBB1 gene for human beta tubulin 1,	1591	100
	3		class VI.	1331	.100
249	gi212834	Gallus gallus	beta-tubulin	1340	85
250	gi13162226	Homo sapiens	Human DNA sequence from clone RP4-543J19 on chromosome 20 Contains part of the GNAS1 gene encoding guanine nucleotide binding protein (G protein, alpha stimulating activity polypeptide 1) including neuroendocrine secretory protein 55 (NESP55), the CTSZA gene encoding cathepsin Z, the ATP5E gene encoding ATP synthase (H+ transporting, mitochondrial F1 complex, epsilon subunit), the gene encoding protein HSPC130 (TH1 Drosophila homolog), the gene for tubulin beta 1 class VI (TUBB1), a gene encoding the CGI-107 protein (LOC51012), four CpG islands, ESTs, STSs and GSSs, complete sequence.	1986	100
250	gil 1230445	Homo sapiens	TUBB1 gene for human beta tubulin 1, class VI.	1986	100
250	gi212834	Galius gallus	beta-tubulin	1699	85
251	gi559325	Homo sapiens	Human mRNA for ATP synthase alpha	1566	99

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
			subunit, complete cds.		
251	gi559317	Homo sapiens	Human gene for ATP synthase alpha subunit, complete cds (exon 1 to 12).	1566	99
251	gi34468	Homo sapiens	H.sapiens mRNA for mitochondrial ATP synthase.	1566	99
252	gi559325	Homo sapiens	Human mRNA for ATP synthase alpha subunit, complete cds.	2192	84
252	gi559317	Homo sapiens	Human gene for ATP synthase alpha subunit, complete cds (exon 1 to 12).	2192	84
252	gi34468	Homo sapiens	H.sapiens mRNA for mitochondrial ATP synthase.	2192	84
253	gi14550508	Homo sapiens	Similar to CG8974 gene product, clone MGC:2460 IMAGE:2964524, mRNA, complete cds.	1051	100
253	gi15928691	Mus musculus	Unknown (protein for MGC:19394)	1036	98
253	gi7293133 .	Drosophila melanogaster	CG8974 gene product	608	66
254	AAE04880	Homo sapiens	Human protease protein-7 (PRTS-7).	2795	100
254	gi14043577	Homo sapiens	hypothetical protein FLJ12455, clone MGC:13149 IMAGE:4298740, mRNA, complete cds.	2795	100
254	AAB94023	Homo sapiens	Human protein sequence SEQ ID NO:14157.	2781	99
255	gi2501855	Homo sapiens	22 kDa actin-binding protein (SM22) gene, complete cds.	937	95
255	gi2340833	Homo sapiens	DNA for SM22 alpha, complete cds.	937	95
255	gi2335047	Homo sapiens	mRNA for SM22 alpha, complete cds.	937	95
256	gi15080204	Homo sapiens	similar to prokaryotic-type class I peptide chain release factors, clone MGC:20261 IMAGE:3029407, mRNA, complete cds.	1948	99
256	gi6706658	Homo sapiens	Human DNA sequence from clone RP1-101K10 on chromosome 6q25-26. Contains a novel gene, the gene for a novel protein similar to Prokaryotic-type class I peptide chain release factors, the 3' end of gene RGS17 (RGSZ2) for regulator of G-protein signaling 17, ESTs, STSs, GSSs and two putative CpG islands, complete sequence.	1940	99
256	gi15680165	Homo sapiens	similar to prokaryotic-type class I peptide chain release factors, clone MGC:20252 IMAGE:4646472, mRNA, complete cds.	1375	98
257	gi15080204	Homo sapiens	similar to prokaryotic-type class I peptide chain release factors, clone MGC:20261 IMAGE:3029407, mRNA, complete cds.	1706	90
257	gi6706658	Homo sapiens	Human DNA sequence from clone RP1-101K10 on chromosome 6q25-26. Contains a novel gene, the gene for a novel protein similar to Prokaryotic-	1698	89

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
			type class I peptide chain release factors, the 3' end of gene RGS17 (RGSZ2) for regulator of G-protein signaling 17, ESTs, STSs, GSSs and two putative CpG islands, complete sequence.		
257	gi15680165	Homo sapiens	similar to prokaryotic-type class I peptide chain release factors, clone MGC:20252 IMAGE:4646472, mRNA, complete cds.	1133	85
258	gi7295482	Drosophila melanogaster	CG4603 gene product	616	41
258	gi12322327	Arabidopsis thaliana	unknown protein	451	46
258	gi9454545	Arabidopsis thaliana	Unknown protein	451	46
259	AAB95307	Homo sapiens	Human protein sequence SEQ ID NO:17548.	5011	100
259	gi14042477	Homo sapiens	cDNA FLJ14740 fis, clone NT2RP3002602, weakly similar to PROBABLE PROTEIN DISULFIDE ISOMERASE ER-60 PRECURSOR (EC 5.3.4.1).	5011	100
259	gi15862252	Homo sapiens	unnamed protein product	5008	99
260	gi15079416	Homo sapiens	secreted modular calcium-binding protein 1, clone MGC:19895 IMAGE:4549051, mRNA, complete cds.	2359	100
260	AAB19394	Homo sapiens	Amino acid sequence of a human secreted protein.	2355	.99
260	gi10432431	Homo sapiens	mRNA for secreted modular calcium- binding protein (smoc1 gene).	2343	.99
261	gi7020475	Homo sapiens	cDNA FLJ20400 fis, clone KAT00587.	1687	100
261	gi1118097	Caenorhabditis elegans	proline and glycine-rich	268	33
261	AAW49723	Homo sapiens	Protein polymer adhesive substrate PPAS1-F.	261	32
262	gi16197949	Drosophila melanogaster	LD21896p	325	29
262	gi7293303	Drosophila melanogaster	CG9089 gene product	325	29
262	gi3170539	Takifugu rubripes	unknown	291	40
263	AAB42525	Homo sapiens	Human ORFX ORF2289 polypeptide sequence SEQ ID NO:4578.	3570	80
263	gi2887497	Homo sapiens	chromosome 19, overlapping cosmids R28707 and R34001, complete sequence.	3570	80
263	AAB42538	Homo sapiens	Human ORFX ORF2302 polypeptide sequence SEQ ID NO:4604.	2835	99
264	gi14017849 .	Homo sapiens	mRNA for KIAA1816 protein, partial cds.	1637	99
264	gi8655687	Homo sapiens	mRNA; cDNA DKFZp762E1511	892	100

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
			(from clone DKFZp762E1511).	1	
264	gi6979930	Homo sapiens	Maml mRNA, partial cds.	315	30
265	gi12836420	Mus musculus	putative	2511	93
265	gi10437002	Homo sapiens	cDNA: FLJ21013 fis, clone CAE05223.	1859	99
265	AAB58385	Homo sapiens	Lung cancer associated polypeptide sequence SEQ ID 723.	1704	99
266	gi14198321	Mus musculus	ribosomal protein L31	543	92
266	gi57115	Rattus norvegicus	ribosomal protein L31 (AA 1-125)	543	92
266	gi14586963	Mus musculus	M75	543	92
267	gi178424	Homo sapiens	Human apolipoprotein A-II mRNA, complete cds.	478	96
267	gi296634	Homo sapiens	Human gene for apolipoprotein AII.	478	96
267	gi296633	Homo sapiens	Human DNA for apolipoprotein A-II.	478	96
268	AAB47184	Homo sapiens	ACPLX protein sequence.	3571	100
268	gi7321168	Homo sapiens	Human DNA sequence from clone RP5-860F19 on chromosome 20p12.3-	3571	100
			13 Contains the gene for KIAA1442 (similar to olfactory neuronal transcription factors (COE1, COE2, COE3, EBF3, OLF1)), RPL19 (60S		
			ribosomal protein L19) and HSPC080 pseudogenes, the gene for metallocarboxypeptidase (CPX-1) and		
٠.			a novel gene. Contains ESTs, STSs, GSSs and four CpG islands, complete sequence.	·	·
268	AAB36174	Homo sapiens	Human APG04 protein.	3567	99
269	gi2314829	Homo sapiens	jerky gene product homolog mRNA, complete cds.	1430	59
269	gi10140857	Mus musculus	jerky	752	33
269	AAG62624	Homo sapiens	Human cell nucleus regulatory protein 56.	598	34
270	gi7959227	Homo sapiens	mRNA for KIAA1483 protein, partial cds.	2231	99
270	gi34192	Homo sapiens	Human KUP mRNA for protein with two zinc fingers.	627	39
270	gi13310782	Mus musculus	myoneurin	315	24
271	AAB93814	Homo sapiens	Human protein sequence SEQ ID NO:13604.	1408	97
271	gi10433080	Homo sapiens	cDNA FLJ11753 fis, clone HEMBA1005583.	1408	97
271	AAB41771	Homo sapiens	Human ORFX ORF1535 polypeptide sequence SEQ ID NO:3070.	821	99
272	gi7959197	Homo sapiens	mRNA for KIAA1468 protein, partial cds.	4603	100
.72	gi15080502	Homo sapiens	clone MGC:16944 IMAGE:4339646, mRNA, complete cds.	4317	94
272	gi9755831	Arabidopsis thaliana	putative protein	675	27
273	gi15080502	Homo sapiens	clone MGC:16944 IMAGE:4339646, mRNA, complete cds.	4362	98

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
273	gi7959197	Homo sapiens	mRNA for KIAA1468 protein, partial cds.	4360	96
273	gi9755831	Arabidopsis thaliana	putative protein	704	28
274	AAB92483	Homo sapiens	Human protein sequence SEQ ID NO:10570.	2626	100
274	gi7021875	Homo sapiens	cDNA FLJ10051 fis, clone HEMBA1001281.	2626	100
274	gi12837616	Mus musculus	putative	2065	90
275	gi10716076	Homo sapiens	mRNA for testis-abundant finger protein, complete cds.	2739	100
275	gi14043332	Homo sapiens	Similar to ring finger protein 23, clone MGC:2475 IMAGE:3051389, mRNA, complete cds.	2533	94
275	gi10716078	Mus musculus	testis-abundant finger protein	2497	92
276	AAB44673	Homo sapiens	Human secreted protein sequence encoded by gene 33 SEQ ID NO:138.	1014	96
276	gi1747	Oryctolagus cuniculus	trichohyalin	213	22
276	gi13936996	Human herpesvirus 8	ORF73	203	22
277	AAG74326	Homo sapiens	Human colon cancer antigen protein SEQ ID NO:5090.	1101	100
277	AAB56461	Homo sapiens	Human prostate cancer antigen protein sequence SEQ ID NO:1039.	778	100
277	gi12842930	Mus musculus	putative	688	90
278	gi1020145	Homo sapiens	Human DNA binding protein (HPF2) mRNA, complete cds.	1528	47
278	gi14456631	Homo sapiens	Human DNA sequence from clone RP1-54B20 on chromosome Xp11.1-11.3. Contains the 5' end of a novel SSX family protein gene, two novel KRAB box containing C2H2 type zinc finger protein genes, a KRAB box protein pseudogene, the gene for a novel protein similar to lysozyme C (1,4-beta-N-acetylmuramidase), the ZNF81 gene for zinc finger protein 81 (HFZ20), ESTs, STSs, GSSs and three CpG islands, complete sequence.	1497	55
278	gi498152	Homo sapiens	Human mRNA for KIAA0065 gene, partial cds.	1495	46
279	gi2914676	Homo sapiens	chromosome 16, cosmid clone 360H6 (LANL), complete sequence.	882	35
279	gi14250678	Homo sapiens	clone MGC:10489 IMAGE:3945548, mRNA, complete cds.	882	35
279	gi2342506	Homo sapiens	mRNA for zinc finger protein FPM315, complete cds.	875	35
280	gi434779	Homo sapiens	Human mRNA for KIAA0112 gene, partial cds.	2072	100
280	gi15278392	Homo sapiens	homolog of yeast ribosome biogenesis regulatory protein RRS1, clone MGC:4831 IMAGE:3603972, mRNA,	1905	100

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
			complete cds.	1	
280 .	gi12804751	Homo sapiens	Similar to regulator for ribosome resistance homolog (S. cerevisiae), clone MGC:2755 IMAGE:2824034, mRNA, complete cds.	1905	100
281	AAB95761	Homo sapiens	Human protein sequence SEQ ID NO:18686.	789	100
281	AAG81272	Homo sapiens	Human AFP protein sequence SEQ ID NO:62.	789	100
281	gi14035852	Homo sapiens	unnamed protein product	789	100
282	gi15080911	Homo sapiens	neo-poly(A) polymerase mRNA, complete cds.	3797	99
282	gi15384858	Homo sapiens	mRNA for poly(A) polymerase gamma (PAPOLG gene).	3797	.99
282	gi13641252	Homo sapiens	SRP RNA 3' adenylating enzyme/pap2 mRNA, complete cds.	3779	99
283	gi6807698	Homo sapiens	mRNA; cDNA DKFZp434A1014 (from clone DKFZp434A1014); partial cds.	1437	85
283	gi12853788	Mus musculus	putative	408	38
283	gi4468790	Xenopus laevis	speedy protein	154	26
284	gi3327062	Homo sapiens	mRNA for KIAA0624 protein, partial cds.	10179	99
284	gi13702612	Staphylococcu s aureus subsp. aureus N315	ORFID:SA2447~hypothetical protein, similar to streptococcal hemagglutinin protein	223	19
284	gi14248429	Staphylococcu s aureus subsp. aureus Mu50	hypothetical protein	223	19
285	gi12697941	Homo sapiens	mRNA for KIAA1698 protein, partial cds.	4716	100
285	gi7299794	Drosophila melanogaster	CG9591 gene product	290	31
285	AAR99256	Homo sapiens	Natural killer lytic associated protein.	92	40
286	AAG62395	Homo sapiens	Human zinc finger protein 46.	2375	100
	gi7576274	Homo sapiens	Human DNA sequence from clone RP11-393J16 on chromosome 10. Contains part of the ZNF33A gene for zinc finger protein 33a (KOX 31), a novel gene for a novel KRAB box containing zinc finger gene, a zinc finger pseudogene, ESTs, STSs, GSSs and two putative CpG islands, complete sequence.	2015	100
286	gi881564	Homo sapiens	Human zinc finger containing protein ZNF157 (ZNF157) mRNA, complete cds.	1339	51
287	gi2822143	Homo sapiens	chromosome 19, cosmid R30217, complete sequence.	1838	53
287	gi9968290	Homo sapiens	mRNA for zinc finger protein (ZNF304 gene).	1735	50
287	gi13543419	Homo sapiens	Similar to zinc finger protein 304,	1735	51

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
,			clone MGC:4079 IMAGE:3530863, mRNA, complete cds.		
288	gi540469	Homo sapiens	(clone HGT26) T cell receptor gamma- chain mRNA, V region.	399	91
288	gi3047024	Homo sapiens	T-cell receptor gamma V1 gene region.	384	100
288	gi339167	Homo sapiens	Human T-cell receptor rearranged gamma-chain gene V-region (V4) (subgroup I).	384	100
289	AAY69976	Homo sapiens	DHFR-HM protein.	886	93
289	gi182724	Homo sapiens	Human dihydrofolate reductase gene.	886	93
289	gi182717	Homo sapiens	Human dihydrofolate reductase gene, exon 6 and 3' flank.	886	93
290	AAE01782	Homo sapiens	Human gene 13 encoded secreted protein HDPNW93, SEQ ID NO:103.	4269	99
290	gi10437433	Homo sapiens	cDNA: FLJ21347 fis, clone COL02724.	4127	97
290	AAB74693	Homo sapiens	Human protease and protease inhibitor PPIM-26.	3948	99
291	gi6681662	Mus musculus	ENH3	955	90
291	gi12844277	Mus musculus	putative	800	79
291	AAY12510	Homo sapiens	Human 5' EST secreted protein SEQ ID NO:541.	648	99
292	AAB47327	Homo sapiens	FCTR4.	2798	98
292	gi15141735	Homo sapiens	unnamed protein product	2798	98
292	gi9663126	Homo sapiens	mRNA for chromosome 12 open reading frame 3 (C12orf3).	214	24
293	gi10440367	Homo sapiens	mRNA for FLJ00018 protein, partial cds.	5938	100
293	gi15488570	Homo sapiens	Similar to hypothetical protein FLJ00018, clone MGC:10073 IMAGE:3896004, mRNA, complete cds.	4736	99
293	gi10438857	Homo sapiens	cDNA: FLJ22458 fis, clone HRC10001.	1570	99
294	AAB08948	Homo sapiens	Human secreted protein sequence encoded by gene 21 SEQ ID NO:105.	1601	99
294	AAB08911	Homo sapiens	Human secreted protein sequence encoded by gene 21 SEQ ID NO:68.	1601	99
294	AAB80238	Homo sapiens	Human PRO238 protein.	641	44
295	AAB18457	Homo sapiens	A human TANGO 216 polypeptide clone.	2106	98
295	AAB18447	Homo sapiens	Amino acid sequence of human TANGO 216 polypeptide.	2106	98
295	gi14017381	Homo sapiens	tumor endothelial marker 8 precursor (TEM8) mRNA, complete cds.	1231	57
296	gi14388342	Macaca fascicularis	hypothetical protein	3833	92
296	gi7243195	Homo sapiens	mRNA for KIAA1407 protein, partial cds.	3817	100
296	gi15451319	Macaca fascicularis	hypothetical protein	2408	91
297	gi7243039	Homo sapiens	mRNA for KIAA1329 protein, partial cds.	4761	100

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
297	gi12007720	Mus musculus	VPS10 domain receptor protein SorCS2	4466	88
297	gi7715916	Mus musculus	SorCSb splice variant of the VPS10 domain receptor SorCS	2177	47
298	AAM00812	Homo sapiens	Human bone marrow protein, SEQ ID NO: 175.	1488	99
298	gi12846045	Mus musculus	putative	1387	65
298	AAM00925	Homo sapiens	Human bone marrow protein, SEQ ID NO: 401.	996	100
299	gi7298852	Drosophila melanogaster	CG10068 gene product	609	43
299	gi8655669	Homo sapiens	mRNA; cDNA DKFZp547C176 (from clone DKFZp547C176).	482	52
299	AAB42048	Homo sapiens	Human ORFX ORF1812 polypeptide sequence SEQ ID NO:3624.	325	46
300	gi14043285	Homo sapiens	Similar to KIAA0808 gene product, clone MGC:15880 IMAGE:3529159, mRNA, complete cds.	1306	97
300	gi7263912	Homo sapiens	Human DNA sequence from clone RP5-1108D11 on chromosome 20q12- 13.11 Contains part of the gene for a novel protein similar to C. elegans	797	96
			T22C1.7, part of the gene for a novel HMG (high mobility group) box protein similar to KIAA0737, KIAA0808 and TNRC9 (CAGF9), ESTs, STSs, GSSs and two putative		
	·	,	CpG islands, complete sequence.		
300	gi3882337	Homo sapiens	mRNA for KIAA0808 protein, complete cds.	767	55
301	gi15430292	Homo sapiens	muscle alpha-kinase (MAK) mRNA, complete cds.	5445	99
301	gi7243041	Homo sapiens	mRNA for KIAA1330 protein, partial cds.	4933	100
301	gi14331137	Mus musculus	myocytic induction/differentiation originator	3684	72
302	gi14550508	Homo sapiens	Similar to CG8974 gene product, clone MGC:2460 IMAGE:2964524, mRNA, complete cds.	589	100
302	gi15928691	Mus musculus	Unknown (protein for MGC:19394)	574	97
302	gi2564951	Mus musculus	unknown	378	72
303	gi7242955	Homo sapiens	mRNA for KIAA1300 protein, partial cds.	9573	99
303	gi6599162	Homo sapiens	mRNA; cDNA DKFZp434N1272 (from clone DKFZp434N1272); partial cds.	1392	98
303	AAG75083	Homo sapiens	Human colon cancer antigen protein SEQ ID NO:5847.	628	92
304	gi1408209	Homo sapiens	Human endogenous retrovirus HERV- K(HML6) proviral clone HML6.17	398	86
304			putative polymerase and envelope genes, partial cds, and 3'LTR.		

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
·		mammary tumor virus			
304	gi6911288	Exogenous mouse mammary tumor virus	Gag-Pro-Pol	176	48
305	gi14269502	Homo sapiens	unconventional myosin 1G valine form (MYO1G) mRNA, MYO1G-V allele, partial cds.	3269	98
305	gi14269504	Homo sapiens	unconventional myosin 1G methonine form (MYO1G) mRNA, MYO1G-M allele, partial cds.	3266	97
305	gi3724141	Rattus norvegicus	myosin I	3130	57
306	gi2145060	Homo sapiens	TTF-I interacting peptide 20 mRNA, partial cds.	2081	99
306	gi2224593	Homo sapiens	Human mRNA for KIAA0326 gene, partial cds.	648	39
306	gi488555	Homo sapiens	Human zinc finger protein ZNF135 mRNA, complete cds.	590	40
307	gi13183883	Homo sapiens	PD-1-ligand 2 protein (PDL2) mRNA, complete cds.	1417	99
307	gi13569410	Homo sapiens	butyrophilin precursor B7-DC mRNA, complete cds.	1417	99
307	AAE01352	Homo sapiens	Human gene 1 encoded secreted protein HDPPA04, SEQ ID NO:74.	1416	99
308	AAB87436	Homo sapiens	Human gene 22 encoded secreted protein fragment, SEQ ID NO:177.	383	100
308	AAB94868	Homo sapiens	Human protein sequence SEQ ID NO:16072.	383	100
308	gi10436314	Homo sapiens	cDNA FLJ13984 fis, clone Y79AA1001846.	383	100
309	AAY85025	Homo sapiens	Human Rap2 amino acid sequence.	206	33
309	gi4678734	Homo sapiens	Human gene from PACs 37M17 and 305B16, chromosome X, similar to small G proteins, especially RAP-2A.	206	33
309	AAM00956	Homo sapiens	Human bone marrow protein, SEQ ID NO: 432.	205	32
310	gi36905	Homo sapiens	Human mRNA for T-cell receptor alpha-chain HAP50 V(a)8.2-J(a)M.	590	100
310	gi1223888	synthetic construct	T cell receptor alpha chain	586	100
310	gi2358036	Homo sapiens	T-cell receptor alpha delta locus from bases 250472 to 501670 (section 2 of 5) of the Complete Nucleotide Sequence.	586	100
311	AAE01596	Homo sapiens	Human gene 13 encoded secreted protein HCLCJ15, SEQ ID NO:146.	1066	92
311	AAE04136	Homo sapiens	Human gene 6 encoded secreted protein HCLBW50, SEQ ID NO:123.	1066	92
311	gi31135	Homo sapiens	H.sapiens mRNA for elongation factor 1-beta.	1066	92
312	gi7243137	Homo sapiens	mRNA for KIAA1378 protein, partial	2400	99

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
	† 		cds.	 	Adendity
312	gi12314036	Homo sapiens	Human DNA sequence from clone RP3-383J4 on chromosome 1q24.1- 24.3 Contains part of a gene encoding a kelch motif containing protein, part of a novel gene encoding a protein similar to Aspartyl-TRNA synthetase, a putative novel gene, a 40S ribosomal protein S27 (RPS27) pseudogene, 2 CpG islands, ESTs, STSs and GSSs,	1184	44
312	gi4650844	Homo sapiens	complete sequence. mRNA for Kelch motif containing protein, complete cds.	1176	44
313	gi7019945	Homo sapiens	cDNA FLJ20079 fis, clone COL03057.	1610	02
313	gi12804721	Homo sapiens	clone MGC:2663 IMAGE:3543910, mRNA, complete cds.	1271	48
313	AAB43912	Homo sapiens	Human cancer associated protein sequence SEQ ID NO:1357.	1255	45
314	AAB41414	Homo sapiens	Human ORFX ORF1178 polypeptide sequence SEQ ID NO:2356.	5094	97
314	gi6329897	Homo sapiens	mRNA for KIAA1137 protein, partial cds.	4798	98
314	gi14043759	Homo sapiens	clone IMAGE:4111596, mRNA, partial cds.	3906	98
315	AAB28375	Homo sapiens	Human hyperpolarisation-activated channel HAC3.	3686	99
315	gi7959337	Homo sapiens	mRNA for KIAA1535 protein, partial cds.	3665	99
315	gi3242244	Mus musculus	hyperpolarization-activated cation channel, HAC3	3556	96
316	gi14198399	Mus musculus	RIKEN cDNA 1500034J20 gene	837	93
316	gi12854536	Mus musculus	putative	837	93
316	gi14250857	Homo sapiens	Human DNA sequence from clone RP5-1137O17 on chromosome 11p12- 14.2 Contains part of a gene similar to putative mitochondrialninner membrane protease subnunit 2, a novel mRNA, ESTs and GSSs, complete sequence.	775	100
317	gi10439850	Homo sapiens	cDNA: FLJ23233 fis, clone CAS00458.	1081	50
317	gi9968290	Homo sapiens	mRNA for zinc finger protein (ZNF304 gene).	1039	48
317	gi14249844	Homo sapiens	Similar to hypothetical protein FLJ23233, clone MGC:14876 IMAGE:3544044, mRNA, complete cds.	1037	47
318	gi11863686	Mus musculus	neurobeachin	3371	96
318	gi11863539	Gallus gallus	neurobeachin	2100	89
318	AAB92596	Homo sapiens	Human protein sequence SEQ ID NO:10843.	1721	100
319	gi12698174	Macaca fascicularis	hypothetical protein	1221	95

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
319	gi10439153	Homo sapiens	cDNA: FLJ22672 fis, clone HSI09265.	1085	99
319	gi7020125	Homo sapiens	cDNA FLJ20190 fis, clone COLF0714.	893	50
320	gi2865219	Homo sapiens	integrin binding protein Del-1 (Del1) mRNA, complete cds.	447.	100
320	AAW94685	Homo sapiens	Human Del-1 protein.	438	98
320	AAW10365	Homo sapiens	Human developmentally-regulated endothelial cell locus-1 protein.	438	98
321	AAB27246	Homo sapiens	Human EXMAD-24 SEQ ID NO: 24.	2047	100
321	AAB42385	Homo sapiens	Human ORFX ORF2149 polypeptide sequence SEQ ID NO:4298.	2047	100
321	gi52998	Mus musculus	macrophage mannose receptor precursor	164	31
322	gi12834087	Mus musculus	putative	1456	82
322	gi2463628	Homo sapiens	Human putative monocarboxylate transporter (MCT) mRNA, complete cds.	506	29
322	gi2198807	Gallus gallus	monocarboxylate transporter 3	473	27
323	gi15620909	Homo sapiens	mRNA for KIAA1925 protein, partial cds.	1059	38
323	AAB92496	Homo sapiens	Human protein sequence SEQ ID NO:10598.	1050	36
323	gi7021900	Homo sapiens	cDNA FLJ10065 fis, clone HEMBA1001455.	1050	36
324	gi9651075	Macaca fascicularis	unnamed protein product	3716	95
324	gi15145795	Sus scrofa	basic proline-rich protein	222	26
324	gi5917666	Zea mays	extensin-like protein	195	25
325	gi7529597	Homo sapiens	Human DNA sequence from clone RP3-402N21 on chromosome 6p21.1- 21.31. Contains up to three novel genes with MAM and immunoglobulin domains. Contains ESTs, STSs, GSSs and four putative CpG islands, complete sequence.	1474	100
325	gi12836077	Mus musculus	putative	1365	95
325	AAE00586	Homo sapiens	Human nuclear cell adhesion molecule homologue, NCAM d 2 protein.	1303	49
326	gi15278193	Homo sapiens	MAGI-1C beta mRNA, complete cds, alternatively spliced.	1492	100
326	gi2702351	Mus musculus	putative membrane-associated guanylate kinase 1	1112	83
326	gi5817255	Homo sapiens	mRNA; cDNA DKFZp434B203 (from clone DKFZp434B203); partial cds.	739	100
327	AAB01432	Homo sapiens	Human TANGO 239 (form 2).	3675	99
327	AAB01426	Homo sapiens	Human TANGO 239.	2700	100
327	AAB00036	Homo sapiens	Human TANGO 239 partial sequence.	2483	97
328	gi7243117	Homo sapiens	mRNA for KIAA1368 protein, partial cds.	5542	100
328	AAY71460	Homo sapiens	Human semaphorin 6A-1.	5422	98
328	gi10187891	Homo sapiens	unnamed protein product	5422	98
329	gi13676461	Macaca fascicularis	hypothetical protein	2193	75

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
329	gi4589566	Homo sapiens	mRNA for KIAA0961 protein, complete cds.	2190	75
329	gi456269	Mus musculus domesticus	zinc finger protein 30	2073	71
330	AAB94295	Homo sapiens	Human protein sequence SEQ ID NO:14747.	3062	99
330	gi10434454	Homo sapiens	cDNA FLJ12768 fis, clone NT2RP2001576, weakly similar to HYPOTHETICAL 62.2 KD PROTEIN C4G8.12C IN CHROMOSOME I.	3062	99
330	gi7291781	Drosophila melanogaster	CG3419 gene product	471	32
331	gi12852801	Mus musculus	putative	1185	95
331	gi12314230	Homo sapiens	Human DNA sequence from clone RP5-846F13 on chromosome 1p21.1- 22.1 Contains part of the PPAP2C (phosphatidic acid phosphatase type 2c) gene, ESTs, STSs and GSSs, complete sequence.	975	100
331	gi7020303	Homo sapiens	cDNA FLJ20300 fis, clone HEP06465.	748	56
332	gi12309630	Homo sapiens	Human DNA sequence from clone RP11-438B23 on chromosome 9 Contains a novel gene for a neuronal leucine-rich repeat protein, ESTs, STSs and GSSs, complete sequence.	3138	100
332	AAB31161	Homo sapiens	Amino acid sequence of a human TOLL protein.	2600	86
332	gi13444976	Homo sapiens	unnamed protein product	2600	86
333	gi4240145	Homo sapiens	mRNA for KIAA0828 protein, partial cds.	3226	99
333	gi14249936	Homo sapiens	Similar to S-adenosylhomocysteine hydrolase-like 1, clone IMAGE:3536052, mRNA, partial cds.	3202	100
333	AAW56097	Homo sapiens	Amino acid sequence of the 0DD4b5.3 enzyme.	2466	84
334	gi13625385	Homo sapiens	EPI64 (EPI64) mRNA, complete cds.	1026	46
334	AAB95321	Homo sapiens	Human protein sequence SEQ ID NO:17577.	1023	50
334	gi10435007	Homo sapiens	cDNA FLJ13130 fis, clone NT2RP3002972, weakly similar to Halocynthia roretzi mRNA for HrPET- 1.	1023	50
335	gi15862408	Homo sapiens	unnamed protein product	2255	95
335	gi13272520	Mus musculus	pancreatitis-induced protein 49	2021	85
335	AAE02778	Homo sapiens	Human PRO-C-MG.64 protein encoded by DNA-C-MG.64-1776 cDNA clone.	1784	95
336	gi15862408	Homo sapiens	unnamed protein product	2281	99
336	gi13272520	Mus musculus	pancreatitis-induced protein 49	2047	88
336	AAE02778	Homo sapiens	Human PRO-C-MG.64 protein encoded by DNA-C-MG.64-1776 cDNA clone.	1810	99
337	gi4545313	Mus musculus	prominin-like protein	1021	77
337	gi15042603	Rattus norvegicus	prominin	647	30

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
337	AAB94028	Homo sapiens	Human protein sequence SEQ ID NO:14170.	642	29
338	gi2978255	Mus musculus	myeloid zinc finger protein-2	212	42
338	AAB54292	Homo sapiens	Human pancreatic cancer antigen protein sequence SEQ ID NO:744.	208	30
338	gi8886436	Homo sapiens	myeloid zinc finger protein 1 splice variants (ZNF42) gene, complete cds, alternatively spliced.	207	42
339	gi3882269	Homo sapiens	mRNA for KIAA0774 protein, partial cds.	5974	99
339	gi12860422	Mus musculus	putative	692	96
339	gi15424451	Homo sapiens	hATIP3	606	36
340	AAB36617	Homo sapiens	Human FLEXHT-39 protein sequence SEQ ID NO:39.	584	100
340	gi8218050	Homo sapiens	Human DNA sequence from clone RP1-187J11 on chromosome 6q11.1-22.33. Contains the gene for a novel protein similar to S. pombe and S. cerevisiae predicted proteins, the gene for a novel protein similar to protein kinase C inhibitors, the 3' end of the gene for a novel protein similar to Drosophila L82 and predicted worm proteins, ESTs, STSs, GSSs and two putative CpG islands, complete sequence.	562	100
340	gi13540300	Mus musculus	nucleolar protein C7B	415	66
341	gi14583268	Homo sapiens	cytoplasmic protein mRNA, complete cds.	628	62
341	gi2104769	Homo sapiens	echinoderm microtubule-associated protein homolog HuEMAP mRNA, complete cds.	560	65
341	gi4406218	Homo sapiens	echinoderm microtubule-associated protein-like EMAP2 mRNA, complete cds.	495	59
342	AAB60099	Homo sapiens	Human transport protein TPPT-19.	1616	93
342	gi7294748	Drosophila melanogaster	CG7616 gene product	580	43
342	gi14714781	Mus musculus	RIKEN cDNA 2610005A10 gene	441	35
343	AAB94374	Homo sapiens	Human protein sequence SEQ ID NO:14915.	3938	99
343	gi10434690	Homo sapiens	cDNA FLJ12921 fis, clone NT2RP2004600.	3938	99
343	gi5689736	Homo sapiens	mRNA for myopodin.	883	34
344	AAY72604	Homo sapiens	Human Electron Transfer Protein, ETRN-2.	717	100
344	gi10953950	Geochelone carbonaria	alpha-D chain hemoglobin	407	54
344	gi4455876	Cairina moschata	alpha D-globin	398	53
345	AAY72604	Homo sapiens	Human Electron Transfer Protein, ETRN-2.	668	78
345	gi10953950	Geochelone	alpha-D chain hemoglobin	359	43

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
		carbonaria			
345	gi4455876	Cairina moschata	alpha D-globin	349	41
346	gi8655669	Homo sapiens	mRNA; cDNA DKFZp547C176 (from clone DKFZp547C176).	1053	100
346	AAB42048	Homo sapiens	Human ORFX ORF1812 polypeptide sequence SEQ ID NO:3624.	840	100
346	gi7298852	Drosophila melanogaster	CG10068 gene product	601	40
347	gi15778899	Homo sapiens	Similar to f-box only protein 17, clone MGC:11162 IMAGE:3841901, mRNA, complete cds.	1537	99
347	gi9280060	Macaca fascicularis	unnamed protein product	1435	95
347	gi15214527	Homo sapiens	Similar to f-box only protein 17, clone MGC:9379 IMAGE:3864760, mRNA, complete cds.	857	56
348	AAG64860	Homo sapiens	Heart muscle cell differentiation related protein SEQ ID NO: 61.	1079	90
348	AAB99931	Homo sapiens	Human MesP1 protein sequence SEQ ID NO:61.	1079	90
348	gi13623241	Homo sapiens	Similar to mesoderm posterior 1, clone MGC:10676 IMAGE:3944350, mRNA, complete cds.	1079	90
349	gi4235144	Homo sapiens	chromosome 19, BAC 39498 (CIT-B-26L23), complete sequence.	387	100
349	gi8163824	Homo sapiens	krueppel-like zinc finger protein HZF2 mRNA, complete cds.	290	74
349	AAY39779	Homo sapiens	CBMACD04 protein sequence.	286	71
350	gi7673618	Mus musculus	ubiquitin specific protease	2016	73
350	gi5689463	Homo sapiens	mRNA for KIAA1063 protein, partial cds.	2000	64
350	gi16198231	Drosophila melanogaster	LD43147p	1188	46
351	gi13540193	Homo sapiens	isopentenyl pyrophosphate isomerase 1 (IDI1), HT009-like protein, and isopentenyl pyrophosphate isomerase type 2 (IDI2) genes, complete cds.	1202	100
351	gi13925766	Homo sapiens	isopentenyl diphosphate dimethylallyl diphosphate isomerase 2 (IDI2) gene, exon 4 and complete cds.	1202	100
351	gi13925769	Homo sapiens	isopentenyl diphosphate dimethylallyl diphosphate isomerase 2 (IDI2) mRNA, complete cds.	1202	100
352	gi13561001	Homo sapiens	Human DNA sequence from clone RP11-528A10 on chromosome 6 Contains an IMPDH1 (IMP (inosine monophosphate) dehydrogenase 1) pseudogene, an RNA helicase pseudogene, a novel gene similar to KIAA0161, ESTs, STSs and GSSs,	950	100
352	gi13991706	Mus musculus	complete sequence. UbcM4-interacting protein 4	655	53

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
352	gi1136384	Homo sapiens	Human mRNA for KIAA0161 gene, complete cds.	651	53
353	gi13561001	Homo sapiens	Human DNA sequence from clone RP11-528A10 on chromosome 6 Contains an IMPDH1 (IMP (inosine monophosphate) dehydrogenase 1) pseudogene, an RNA helicase	709	79
			pseudogene, a novel gene similar to KIAA0161, ESTs, STSs and GSSs, complete sequence.		
353	gi13991706	Mus musculus	UbcM4-interacting protein 4	506	45
353	gi1136384	Homo sapiens	Human mRNA for KIAA0161 gene, complete cds.	502	44
354	AAB74446	Homo sapiens	Human protease-inhibitor like protein.	2759	100
354	gi12053227	Homo sapiens	mRNA; cDNA DKFZp434B044 (from clone DKFZp434B044); complete cds.	2756	99
354	gi15593902	Homo sapiens	unnamed protein product	2743	99
355	AAB94358	Homo sapiens	Human protein sequence SEQ ID NO:14883.	1788	98
355	gi10434632	Homo sapiens	cDNA FLJ12886 fis, clone NT2RP2004041, weakly similar to SYNAPSINS IA AND IB.	1788	98
355	gi12052738	Homo sapiens	mRNA; cDNA DKFZp564H1322 (from clone DKFZp564H1322); complete cds.	1788	98
356	gi13436437	Homo sapiens	Similar to RIKEN cDNA 5730438N18 gene, clone MGC:4399 IMAGE:2905957, mRNA, complete cds.	1634	99
356	gi15030091	Mus musculus	Similar to RIKEN cDNA 5730438N18 gene	1508	91
356	AAB43372	Homo sapiens	Human ORFX ORF3136 polypeptide sequence SEQ ID NO:6272.	1464	91
357	AAB73511	Homo sapiens	Human transferase HTFS-18, SEQ ID NO:18.	1880	99
357	AAG74560	Homo sapiens	Human colon cancer antigen protein SEQ ID NO:5324.	450	98
357	AAG02792	Homo sapiens	Human secreted protein, SEQ ID NO: 6873.	324	96
358	gi7673618	Mus musculus	ubiquitin specific protease	2711	95
358	gi5689463	Homo sapiens	mRNA for KIAA1063 protein, partial cds.	2382	78
358	gi5823525	Drosophila melanogaster	ubiquitin-specific protease nonstop	1305	49
359	AAB94775	Homo sapiens	Human protein sequence SEQ ID NO:15864.	1022	100
359	gi10435984	Homo sapiens	cDNA FLJ13842 fis, clone THYRO1000793.	1022	100
359	gi2340162	Xenopus laevis	dsRBP-ZFa	380	44
360	gi3676086	bacteriophage PS119	gp19	291	59
360	gi1778468	Escherichia	hypothetical protein	287	59

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
		coli			1
360	gi1786768	Escherichia coli K12	bacteriophage lambda lysozyme homolog	287	59
361	gi13544003	Homo sapiens	clone IMAGE:3677165, mRNA, partial cds.	2172	88
361	gi3169073	Schizosacchar omyces pombe	phenylalanyl-trna synthetase, mitochondrial precursor	233	33
361	gi13877969	Arabidopsis thaliana	putative phenylalanine-tRNA synthetase	228	35
362	gi293694	Mus musculus	laminin receptor	370	49
362	gi13277921	Mus musculus	laminin receptor 1 (67kD, ribosomal protein SA)	367	49
362	gi4633839	Mus musculus	37kDa oncofetal antigen	367	49
363	gi15082271	Homo sapiens	testes development-related NYD-SP21 mRNA, complete cds.	1876	100
363	gi6807923	Homo sapiens	mRNA; cDNA DKFZp434H092 (from clone DKFZp434H092); partial cds.	1620	100
363	gi7294427	Drosophila melanogaster	CG8797 gene product	118	21
364	AAE01355	Homo sapiens	Human gene 4 encoded secreted protein HRABV43, SEQ ID NO:77.	2724	97
364	gi12836042	Mus musculus	putative	2607	93
364	AAE01380	Homo sapiens	Human gene 4 encoded secreted protein HRABV43, SEQ ID NO:102.	2500	97
365	gi10439688	Homo sapiens	cDNA: FLJ23109 fis, clone LNG07754.	2809	99
365	gi9622093	Mus musculus	E-cadherin binding protein E7	2768	97
365	AAG01765	Homo sapiens	Human secreted protein, SEQ ID NO: 5846.	737	99
366 .	gi12854995	Mus musculus	putative	844	71
366	gi10241691	Homo sapiens	Novel human gene mapping to chomosome 22.	791	99
366	gi14602790	Homo sapiens	DKFZP566F0546 protein, clone MGC:2444 IMAGE:2822570, mRNA, complete cds.	791	99
367	gi15082283	Homo sapiens	Similar to small glutamine-rich tetratricopeptide repeat (TPR)- containing, clone MGC:10496 IMAGE:3625993, mRNA, complete cds.	720	100
367	gi3377591	Homo sapiens	full length insert cDNA YN88E09.	592	100
367	gi15488015	Homo sapiens	TPR-containing co-chaperone mRNA, complete cds.	450	64
368	gi9104819	Xylella fastidiosa 9a5c	hypothetical protein	151	43
368	AAY59981	Homo sapiens	Human endometrium turnour EST encoded protein 41.	128	46
368	AAE03351	Homo sapiens	Human gene 4 encoded secreted protein fragment, SEQ ID NO:126.	121	58
369	gi5817053	Homo sapiens	mRNA; cDNA DKFZp586D0824 (from clone DKFZp586D0824); partial cds.	571	43
369	gi15530285	Homo sapiens	clone MGC:24275 IMAGE:3950542,	571	43

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
			mRNA, complete cds.		
369	gi13569476	Mus musculus	immunity-associated nucleotide 4	540	42
370	gi8453103	Homo sapiens	zinc finger protein mRNA, complete cds.	1296	58
370	gi15012179	Homo sapiens	zinc finger protein 16 (KOX 9), clone MGC:15145 IMAGE:3949487, mRNA, complete cds.	1296	58
370	gi498721	Homo sapiens	H.sapiens HZF10 mRNA for zinc finger protein.	1279	55
371	gi15929964	Homo sapiens	Similar to hypothetical protein FLJ10702, clone MGC:21954 IMAGE:4391821, mRNA, complete cds.	973	100
371	AAB42336	Homo sapiens	Human ORFX ORF2100 polypeptide sequence SEQ ID NO:4200.	932	93
371	AAB93080	Homo sapiens	Human protein sequence SEQ ID NO:11912.	923	91
372	gi7328451	Mus musculus	sialidase	893	44
372	AAB93971	Homo sapiens	Human protein sequence SEQ ID NO:14038.	866	42
372	AAW73964	Homo sapiens	Human sialidase protein sequence.	866	42
373	gi1480005	Mus musculus	Zic4 protein	1490	86
373	AAB14349	Homo sapiens	Human Zic1 protein.	1102	67
373	gi1208429	Homo sapiens	mRNA for Zic protein, complete cds.	1102	67
374	gi12860114	Mus musculus	putative	876	40
374	gi161958	Trypanosoma cruzi	surface antigen	177	23
374	gi1334643	Xenopus laevis	APEG precursor protein	174	26
375	AA¥99349	Homo sapiens	Human PRO1110 (UNQ553) amino acid sequence SEQ ID NO:31.	1683	100
375	AAB19729	Homo sapiens	Human SECX Clone 4339264-2 encoded protein.	1683	100
375	AAB15549	Homo sapiens	Human immune system molecule from Incyte clone 2774913.	1683	100
376	gi12746394	Homo sapiens	CUG-BP and ETR-3 like factor 4 (CELF4) mRNA, complete cds.	936	100
376	gi13278792	Homo sapiens	Bruno (Drosophila) -like 4, RNA binding protein, clone MGC:2693 IMAGE:2820541, mRNA, complete cds.	911	98
376	gi12804985	Homo sapiens	Similar to etr1, clone MGC:4320 IMAGE:2820541, mRNA, complete cds.	911	98
377	gi12746394	Homo sapiens	CUG-BP and ETR-3 like factor 4 (CELF4) mRNA, complete cds.	905	89
377	gi13278792	Homo sapiens	Bruno (Drosophila) -like 4, RNA binding protein, clone MGC:2693 IMAGE:2820541, mRNA, complete cds.		88
377	gi12804985	Homo sapiens	Similar to etr1, clone MGC:4320 IMAGE:2820541, mRNA, complete cds.	880	88

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	%
378	gi12841060	Mus musculus	putative	809	Identity 75
378	gi7293285	Drosophila melanogaster	CG4768 gene product	239	37
378	gi1938566	Caenorhabditis elegans	Hypothetical protein C48B6.3	123	38
379	gi3880385	Caenorhabditis elegans	predicted using Genefinder-contains similarity to Pfam domain: PF01484 (Nematode cuticle collagen N-terminal domain), Score=51.5, E-value=6.1e-12, N=1-cDNA EST yk94a4.5 comes from this gene-cDNA EST yk94a4.3 comes from this gene-cDNA EST yk68d1.5 comes from this gene-cDNA EST yk68d1.3 comes from this gene-cDNA EST yk68d1.3 comes from this gene-	79	35
379	gi6684	Caenorhabditis elegans	unnamed protein product	79	35
379	gi156262	Caenorhabditis elegans	collagen	79	35
380	AAB85365	Homo sapiens	Novel Von Willebrand/thrombosporin- like mature protein sequence.	657	94
380	AAB85364	Homo sapiens	Novel Von Willebrand/thrombosporin- like polypeptide.	657	94
380	gi12836633	Mus musculus	putative	651	59
381	gi15024264	Mus musculus	ribosomal protein L35a	191	53
381	gi57119	Rattus norvegicus	ribosomal protein L35a (aa 1-110)	191	53
381	gi12846322	Mus musculus	putative	191	53
382	gi12835133	Mus musculus	putative	617	71
382	gi7293113	Drosophila melanogaster	CG12379 gene product	283	72
382	gi6042159	Caenorhabditis elegans	Hypothetical protein F53A3.7	226	55
383	AAB81053	Homo sapiens	Human protein HP01640 amino acid sequence.	932	100
383	gi12841896	Mus musculus	putative	925	98
383	gi7303144	Drosophila melanogaster	CG10153 gene product	612	65
384	gi10440373	Homo sapiens	mRNA for FLJ00022 protein, partial cds.	1345	93
384	gi10440396	Homo sapiens	mRNA for FLJ00031 protein, partial cds.	647	88
384	gi1086626	Caenorhabditis elegans	Hypothetical protein C06A6.3	273	33
385	gi12053305	Homo sapiens	mRNA; cDNA DKFZp434G099 (from clone DKFZp434G099); complete cds.	1210	100
385	gi2516239	Mus musculus	Rab33B	1138	94
385	gi12836564	Mus musculus	putative	1138	94
386	gi7243247	Homo sapiens	mRNA for KIAA1433 protein, partial cds.	3232	100
386	AAB94053	Homo sapiens	Human protein sequence SEQ ID NO:14222.	3223	99
386	gi13096872	Mus musculus	Unknown (protein for MGC:7720)	2906	89
387	gi14599491	Homo sapiens	small proline-rich protein 2F (SPRR2F)	458	100

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
			gene, complete cds.		
387	gi14599489	Homo sapiens	small proline-rich protein 2E (SPRR2E) gene, complete cds.	444	95
387	gi338423	Homo sapiens	Human small proline rich protein (sprII) mRNA, clone 930.	434	94
388	gi6010699	Rattus norvegicus	F-box protein FBL2	1449	99
388	gi14043139	Homo sapiens	RIKEN cDNA 2610511F20 gene, clone MGC:15482 IMAGE:2987858, mRNA, complete cds.	1383	100
388	gi12848653	Mus musculus	putative	1371	99
389	gi2853265	Rattus norvegicus	jun dimerization protein 2	800	96
389	gi12248392	Mus musculus	transcriptional inhibitory factor	795	95
389	gi6648146	Homo sapiens	chromosome 14 clone CTD-2317F5 map 14q24.3, complete sequence.	481	100
390	gi15277240	Homo sapiens	genomic DNA, chromosome 6p21.3, HLA Class I region, section 17/20.	1296	100
390	gi11875405	Homo sapiens	HZFw1 protein mRNA, complete cds.	1291	99
390	gi11875407	Homo sapiens	HZFw2 protein mRNA, complete cds.	773	99
391	gi6572201 gi4469186	Homo sapiens Homo sapiens	ens Human DNA sequence from clone CITF22-27C3 on chromosome 22q13.1-13.31 Contains a gene for a novel protein (DJ1163J1.2) and part of a gene for a novel protein (DJ1163J1.3, similar to mouse B99), ESTs, STSs and GSSs, complete sequence.		100
201	AAD02551		RP5-1163J1 on chromosome 22q13.2-13.33 Contains the 3' part of a gene for a novel KIAA0279 LIKE EGF-like domain containing protein (similar to mouse Celsr1, rat MEGF2), a novel gene for a protein similar to C. elegans B0035.16 and bacterial tRNA (5-Methylaminomethyl-2-thiouridylate)-Methyltransferases, and the 3' part of a novel gene for a protein similar to mouse B99. Contains ESTs, GSSs and putative CpG islands, complete sequence.		
391	AAB92551	Homo sapiens			96
392	gi5001720	Mus musculus	odd-skipped related 1 protein	1413	97
392	gi15778246	Mus musculus	odd-skipped related 2	924	66
392	gi15488723	Mus musculus	S Unknown (protein for MGC:19171) 924 6		66
393	AAB94364	Homo sapiens			99
393	gi10434650	Homo sapiens			99
393	gi13623217	Homo sapiens	Similar to hypothetical protein FLJ12895, clone IMAGE:3533093,	2150	99

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
			mRNA, partial cds.		1
394	gi12053105	Homo sapiens	mRNA; cDNA DKFZp434K111 (from clone DKFZp434K111); complete cds.	3116	100
394	gi2282582	Mus musculus	actin-binding protein	2402	74
394	AAR94386	Homo sapiens	Human neural cell protein marker RR/B.	2400	74
395	gi207145	Rattus norvegicus	synaptotagmin II	2128	95
395	gi7739733	Mus musculus	synaptotagmin II	2121	95
395	gi688412	Mus musculus	synaptotagminII/IP4BP	2121	95
396	gi15487674	Homo sapiens	OSBP-related protein 1 mRNA, complete cds.	3220	99
396	AAB92611	Homo sapiens	Human protein sequence SEQ ID NO:10880.	703	100
396	AAY97291	Homo sapiens	Lipid associated protein (LIPAP) 2764333CD1.	703	100
397	gi11231085	Macaca fascicularis	hypothetical protein	490	76
397	gi2447128	Paramecium bursaria Chlorella virus 1	contains 10 ankyrin-like repeats; similar to human ankyrin, corresponds to Swiss-Prot Accession Number P16157	212	33
397	gi6634025	Homo sapiens	mRNA for KIAA0379 protein, partial cds.	203	38
398	AAB21047	Homo sapiens	Human nucleic acid-binding protein, NuABP-51.	1082	100
398	gi833629	Xenopus laevis	nucleoplasmin	459	49
398 ·	gi64940	Xenopus laevis	nucleoplasmin (AA 1-200)	435	46
399	gi15919272	Homo sapiens	putative forkhead/winged-helix transcription factor (FOXP2) mRNA, complete cds.	596	84
399	gi2565057	Homo sapiens	CAGH44 mRNA, partial cds.	596	84
399	gi14582802	Mus musculus	forkhead-related transcription factor 2	588	82
400	AAB08199	Homo sapiens	Amino acid sequence of human diacylglycerol kinase beta (DAGKbeta).	4217	99
400	gi10279722	Homo sapiens	unnamed protein product	4217	99
400	gi485398	Rattus norvegicus	90kDa-diacylglycerol kinase	4046	95
401	gi7670446	Mus musculus	unnamed protein product	1295	87
401	gi13185203	Homo sapiens	unnamed protein product	799	83
401	AAY31642	Homo sapiens	Human transport-associated protein-4 (TRANP-4).		35
402	gi12837990	Mus musculus	putative	985	69
402	gi5668737	Mus musculus	UBE-1c2	661	50
402	AAB94645	Homo sapiens	s Human protein sequence SEQ ID 426 NO:15538.		52
403	gi10439821	Homo sapiens	cDNA: FLJ23209 fis, clone ADSH00512.	2596	99
403	gi10440353	Homo sapiens	mRNA for FLJ00011 protein, partial	1448	97

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
			cds.	 	Idenat
403	gi8217420	Homo sapiens	Human DNA sequence from clone	1026	100
	1		RP11-108L7 on chromosome 10.		
		1	contains part of the gene for a novel		1.
	1		Insulin-like growth factor binding type	1	1
	1		protein with Kazal-type serine protease	- [1
		İ	inhibitor domain, the gene for a novel	1	
	1		protein similar to rat tricarboxylate	ŀ	1
			carrier, the gene for a novel PDZ		
			(DHR, GLGF) domain protein, the	1	}
] .	gene for a novel protein similar to	1]
			KIAA0552, KIAA0341 and Fugu	1	
•			hypothetical protein 2, the gene for a		l
	<u> </u>		novel protein similar to Plasmodium	1	
			POM1 and C. elegans F46G11.1, a	l	ļ
•			putative novel gene, the SEMA4G gene		1
		1	for semaphorin 4G and a novel gene.		1
	1	1	Contains ESTs, STSs, GSSs and seven]
			putative CpG islands, complete	1	
			sequence.		
404	AAB42219	Homo sapiens	Human ORFX ORF1983 polypeptide	2230	96
 	<u> </u>		sequence SEQ ID NO:3966.		-0
404	gi3417297	Homo sapiens	Human Chromosome 16 BAC clone	2230	96
			CIT987SK-A-635H12, complete	1	
			sequence.	1	
404	gi15559282	Homo sapiens	clone MGC:20208 IMAGE:3936339,	1021	53
			mRNA, complete cds.		,50
405	gi13365905	Macaca	hypothetical protein	1154	99
405		fascicularis			
405	AAB15537	Homo sapiens	Human immune system molecule from	911	100
105	1.170.000		Incyte clone 2751129.		
405	AAE04891	Homo sapiens	Human transporter and ion channel-4	360	39
406	-:262042	 	(TRICH-4) protein.		
406	gi262843	Rattus sp.	neurotransmitter transporter	3709	96
100	gi545078	Rattus sp.	Na+/Cl(-)-dependent neurotransmitter	3694	96
406	A A D 00000		transporter		
100	AAR88390	Homo sapiens	Human neurotransmitter transporter	3668	96
107	AAB31212	T7	protein.		
,	WD31217	Homo sapiens	Amino acid sequence of human	728	100
107	A A D 44221	YY	polypeptide PRO6004.		
107	AAB44331	Homo sapiens	Human PRO4993 protein sequence	717	100
107	~i4510560	D.#	SEQ ID NO:612.		
.	gi4519558	Rattus	Kilon	667	94
108	ci15277072	norvegicus	G: 11		
100	gi15277972	Mus musculus	Similar to DnaJ (Hsp40) homolog,	808	49
108	ci7804472	Maria	subfamily B, member 1		
108	gi7804472	Mus musculus	heat shock protein 40	808	49
09	AAB72675	Homo sapiens	Human HDJ1.	804	48
.09	gi12841015	Mus musculus	putative	798	52
09	AAB60114	Homo sapiens	Human transport protein TPPT-34.	787	51
עטי	gi13435410	Mus musculus	Similar to RIKEN cDNA 1810012H11	768	53
10	1400555		gene		
10	gi488555	Homo sapiens	Human zinc finger protein ZNF135	1241	52

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
			mRNA, complete cds.		
410	AAY73346	Homo sapiens	HTRM clone 619699 protein sequence.	1238	49
410	AAB43912	Homo sapiens	Human cancer associated protein sequence SEQ ID NO:1357.	1231	49
411	gi837292	Rattus norvegicus	S100A1 gene product	278	59
411	AAB45531	Homo sapiens	Human S100A1 protein.	274	57
411	gi11228039	Homo sapiens	S100A1 cDNA	274	57
412	AAB19851	Homo sapiens	Human muscle-specific protein Ozz.	1504	100
412	gi13929456	Homo sapiens	Human DNA sequence from clone RP3-337018 on chromosome 20q12- 13.1. Contains the PLPT gene encoding Phospholipid Transfer Protein, the PPGB gene coding for Lysosomal Protective Protein precursor (EC 3.4.16.5, Cathepsin A,	1504	100
			Carboxypeptidase C) and the gene encoding peroxisomal acyl-CoA thioesterase (PTE1, thioesterase II), four novel genes, the gene for a novel protein similar to Drosophila Neuralized (Neu) and the 5' end of an isoform of the TNNC2 gene for fast troponin C2. Contains three CpG islands, ESTs, STSs and GSSs, complete sequence.		
412	gi12835750	Mus musculus	putative	1328	89
413	gi12847182	Mus musculus	putative	875	87
413	gi4884173	Homo sapiens	mRNA; cDNA DKFZp564G0982 (from clone DKFZp564G0982); partial cds.	646	100
413	gi10047333	Homo sapiens	mRNA for KIAA1628 protein, partial cds.	346	42
414	gi7959343	Homo sapiens	mRNA for KIAA1538 protein, partial cds.	3286	100
414	AAB42721	Homo sapiens	Human ORFX ORF2485 polypeptide sequence SEQ ID NO:4970.	382	100
414	AAB42764	Homo sapiens	Human ORFX ORF2528 polypeptide sequence SEQ ID NO:5056.	355	41
415	gi14043332	Homo sapiens	Similar to ring finger protein 23, clone MGC:2475 IMAGE:3051389, mRNA, complete cds.	1006	43
415	gi10716078	Mus musculus	testis-abundant finger protein	995	42
415	gi10716076	Homo sapiens	mRNA for testis-abundant finger protein, complete cds.	966	40
416	gi3599509	Mus musculus	rho/rac-interacting citron kinase	1507	61
416	gi3360512	Rattus norvegicus	Citron-K kinase 150		89
416	gi3599507	Mus musculus	rho/rac-interacting citron kinase short 1503 isoform		89
417	gi2358070	Mus musculus	trypsinogen 1	898	65
417	gi603903	Gallus gallus	trypsinogen	408	36
	gi65163	Xenopus	# Thornoxcii	400 .	20

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
·		laevis		7	1
418	gi440127	Rattus norvegicus	cerebroglycan	1132	87
418	AAB44256	Homo sapiens	Human PRO705 (UNQ369) protein sequence SEQ ID NO:109.	570	46
418	AAY25909	Homo sapiens	Human GPC6 protein.	570	46
419	AAM06489	Homo sapiens	Human foetal protein, SEQ ID NO: 220.	376	82
419	gi12835376	Mus musculus	putative	230	31
419	AAE02058	Homo sapiens	Human four disulfide core domain (FDCD)-containing protein.	222	31
420	AAB42561	Homo sapiens	Human ORFX ORF2325 polypeptide sequence SEQ ID NO:4650.	5075	100
420	gi5419865	Homo sapiens	mRNA; cDNA DKFZp434N074 (from clone DKFZp434N074).	5070	99
420	gi4589532	Homo sapiens	mRNA for KIAA0944 protein, partial cds.	3375	61
421	gi10438804	Homo sapiens	cDNA: FLJ22419 fis, clone HRC08593.	1026	60
421	gi13938187	Homo sapiens	hypothetical protein FLJ22419, clone MGC:14900 IMAGE:3347783, mRNA, complete cds.	1026	60
421	gi6690339	Mus musculus	hematopoietic zinc finger protein	717	47
422	AAB94721	Homo sapiens	Human protein sequence SEQ ID NO:15739.	1678	99
422	gi10435784	Homo sapiens	cDNA FLJ13693 fis, clone PLACE2000111.	1678	99
422	gi5706454	Homo sapiens	mRNA for Natural killer cell p44 related gene 2 (NKp44RG2).	158	29
423	gi15026974	Homo sapiens	mRNA for obscurin (OBSCN gene).	2713	96
423	AAB95162	Homo sapiens	Human protein sequence SEQ ID NO:17205.	1173	86
423	gi13938170	Homo sapiens	clone IMAGE:2961284, mRNA, partial cds.	540	26
424	gi12861364	Mus musculus	putative	523	51
424	AAE02058	Homo sapiens	Human four disulfide core domain (FDCD)-containing protein.	485	38
424	gi12655452	Homo sapiens	mRNA for keratin associated protein 4.7 (KRTAP4.7 gene).	485	40
425	gi12830335	Homo sapiens	Human DNA sequence from clone RP11-550O8 on chromosome 20. Contains a novel gene encoding a protein kinase, an RPL7 (60S Ribosomal Protein L7) pseudogene, a CpG island, ESTs, STSs and GSSs, complete sequence.	2062	99
425	AAB65688	Homo sapiens	Novel protein kinase, SEQ ID NO: 216.	1732	100
425	AAB65690	Homo sapiens	Novel protein kinase, SEQ ID NO: 218.		69
426	gi388518	Homo sapiens			95
426	gi36173	Homo sapiens	H.sapiens rearranged T-cell receptor beta chain mRNA.	613	94
426	gi1552509	Homo sapiens	Human germline T-cell receptor beta	606	100

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
			chain TCRBV13S1, TCRBV6S8A2T, TCRBV5S6A3N2T, TCRBV13S6A2T, TCRBV6S9P, TCRBV5S3A2T, TCRBV13S8P, TCRBV6S3A1N1T, TCRBV5S2, TCRBV6S6A2T, TCRBV5S7P, TCRBV13S4,		
			TCRBV6S2A1N1T, TCRBV5S4A2T, TCRBV6S4A1, TCRBV23S1A2T, TCRBV12S1A1N2, TCRBV21S2A2, TCRBV8S1, TCRBV8S2A1T, TCRBV8S3, TCRBV16S1A1N1, TCRBV24S1A3T, TCRBV25S1A2PT, TCRBV26S1P, TCRBV18S1, TCRBV17S1A1T, TCRBV2S1, TCRBV10S1P genes from bases		
40.5			257519 to 472940 (section 2 of 3).		
427	AAE04752	Homo sapiens	Human beta-1,3-galactosyltransferase homologue, ZNSSP8.	434	33
427	gi14597533	Homo sapiens	unnamed protein product	434	33
427	gi14039836	Homo sapiens	beta 1,3 N- acetyglucosaminyltransferase Lc3 synthase mRNA, complete cds.	434	33
428	gi596142	Homo sapiens	Human proteasome subunit LMP7 (allele LMP7C) mRNA, complete cds.	628	49
428	gi38482	Homo sapiens	H.sapiens gene for major histocompatibility complex encoded proteasome subunit LMP7.	624	49
428	gi1054747	Homo sapiens	H.sapiens DMA, DMB, HLA-Z1, IPP2, LMP2, TAP1, LMP7, TAP2, DOB, DQB2 and RING8, 9, 13 and 14 genes.	624	49.
429	AAG71415	Homo sapiens	Human olfactory receptor polypeptide, SEQ ID NO: 1096.	1587	100
429	AAG71594	Homo sapiens	Human olfactory receptor polypeptide, SEQ ID NO: 1275.	1344	83
429	AAG72476	Homo sapiens	Human OR-like polypeptide query sequence, SEQ ID NO: 2157.	1011	100
430	gi10440063	Homo sapiens	cDNA: FLJ23392 fis, clone HEP17418.	3045	100
430	gi15214571	Mus musculus	Unknown (protein for IMAGE:4207025)	2396	80
430	gi1770528	Homo sapiens	H.sapiens mRNA for translin associated zinc finger protein-1.	· 687	38
431	gi12859929	Mus musculus	putative	917	96
431	gi15207935	Macaca fascicularis	hypothetical protein	301	96
431	gi1655637	Mus musculus	orf	147	27
432	gi4585414	Bacteriophage 933W	hypothetical protein	408	42
432	gi4499798	Bacteriophage 933W	orf15; homologous to ninG gene	408	42
432	gi5881629	Bacteriophage VT2-Sa	ge hypothetical protein 408		42
433	gi13161184	Homo sapiens	cytochrome P450 2S1 (CYP2S1) mRNA, complete cds.	2615	100

Table 2

SEQ ID NO:	Accession No.	Species	Description	Score	% Identity
433	AAB93056	Homo sapiens	Human protein sequence SEQ ID NO:11860.	2527	100
433	gi14042396	Homo sapiens	cDNA FLJ14699 fis, clone NT2RP2006571, moderately similar to CYTOCHROME P450 2G1 (EC 1.14.14.1).	2527	100
434	gi13445575	Homo sapiens	facilitative glucose transporter GLUT10 (SLC2A10) mRNA, complete cds.	2752	99
434	gi13603727	Homo sapiens	glucose transporter (GLUT10) mRNA, complete cds.	2752	99
434	gi11065680	Homo sapiens	Novel human gene mapping to chromosome 20, similar to membrane transporters.	2752	99
435	gi13310486	Homo sapiens	C2H2 zinc finger protein (SALL3) gene, complete cds.	6094	99
435	gi6688241	Homo sapiens	SALL3 gene, exons 1a, 2 and 3.	6070	99
435	gi1296845	Mus musculus	spalt protein	5089	84
436	AAG71445	Homo sapiens	Human olfactory receptor polypeptide, SEQ ID NO: 1126.	1312	85
436	AAG71447	Homo sapiens	Human olfactory receptor polypeptide, SEQ ID NO: 1128.	924	61
436	gi15293797	Homo sapiens	clone OR6M1 olfactory receptor gene, partial cds.	829	78
437	AAB65297	Homo sapiens	Human PRO9828 protein sequence SEQ ID NO:511.	1360	100
437	AAG89178	Homo sapiens	Human secreted protein, SEQ ID NO: 298.	1360	100
437	AAB84652	Homo sapiens	Amino acid sequence of fibroblast growth factor homologue zFGF12.	1360	100
438	gi53756	Mus musculus	minopontin precursor (AA -66 to 272)	1521	100
438	gi297546	Mus musculus	osteopontin 1516		99
438	gi50864	Mus musculus	T lymphocyte activation protein	1514	99

Table 3

SEQ ID NO:	Database entry ID	Description	*Results
1	PF00204	Zinc finger C-x8-C-x5-C-x3-H type (and similar).	PF00204 11.59 9.700e-12 426-437
1	BL00518	Zinc finger, C3HC4 type (RING finger), proteins.	BL00518 12.23 3.667e-09 33-42
2	BL00291	Prion protein.	BL00291A 4.49 8.759e-09 185-220
3	PF01105	emp24/gp25L/p24 family.	PF01105B 25.12 1.000e-40 178-230
4	BL00307	Legume lectins beta-chain proteins.	BL00307G 9.91 8.531e-10 678-689
4	PF00922	Vesiculovirus phosphoprotein.	PF00922A 19.17 8.862e-09 281-315
6	BL01159	WW/rsp5/WWP domain proteins.	BL01159 13.85 6.073e-09 61-76
6	BL00591	Glycosyl hydrolases family 10 proteins.	BL00591G 9.65 9.167e-09 311-323
7	BL01159	WW/rsp5/WWP domain proteins.	BL01159 13.85 6.073e-09 61-76
7	BL00591	Glycosyl hydrolases family 10 proteins.	BL00591G 9.65 9.167e-09 311-323
9	BL00913	Iron-containing alcohol dehydrogenases	BL00913D 24.20 8.981e-17 170-204
	ł	proteins.	BL00913C 7.62 4.375e-11 136-146
		•	BL00913B 10.94 7.706e-11 86-102
10	BL00913	Iron-containing alcohol dehydrogenases	BL00913D 24.20 8.981e-17 218-252
		proteins.	BL00913C 7.62 4.375e-11 184-194
			BL00913B 10.94 7.706e-11 134-150
11	BL50062	BCL2-like apoptosis inhibitors (spans part of BH3, BH1 and BH.	BL50062C 6.66 8.500e-11 349-358
14	BL01144	Ribosomal protein L31e proteins.	BL01144 25.07 9.069e-26 78-130
15	PF00204	Zinc finger C-x8-C-x5-C-x3-H type (and similar).	PF00204 11.59 6.694e-10 355-366
15	BL00904	Protein prenyltransferases alpha subunit repeat proteins proteins.	BL00904A 8.30 4.000e-09 485-535
15	BL00415	Synapsins proteins.	BL00415N 4.29 6.727e-12 483-527
	·		BL00415N 4.29 2.774e-09 118-600
			BL00415P 2.37 4.290e-09 819-855
			BL00415Q 2.23 6.534e-09 474-510
15	PR00049	WILM'S TUMOUR PROTEIN	PR00049D 0.00 4.500e-14 490-505
		SIGNATURE	PR00049D 0.00 2.500e-12 489-504
]		PR00049D 0.00 4.000e-12 491-506
	,	•	PR00049D 0.00 8.201e-11 488-503
]		PR00049D 0.00 1.205e-10 492-507
			PR00049D 0.00 3.746e-09 487-502
			PR00049D 0.00 5.271e-09 485-500
			PR00049D 0.00 6.644e-09 493-508
15 .	DM00215	PROLINE-RICH PROTEIN 3.	DM00215 19.43 9.022e-13 471-504
	ľ	•	DM00215 19.43 1.458e-09 483-516
		•	DM00215 19.43 2.678e-09 469-502
			DM00215 19.43 5.424e-09 468-501
		•	DM00215 19.43 8.017e-09 470-503
		·	DM00215 19.43 9.085e-09 466-499
15	BL01113	Cla domain protein-	DM00215 19.43 9.237e-09 484-517
15	BL00048	C1q domain proteins. Protamine P1 proteins.	BL01113A 17.99 9.308e-09 116-143
17		Frommie Fr proteins.	BL00048 6.39 5.263e-10 196-223 BL00048
	•		6.39 3.363e-09 262-289 BL00048 6.39
17	PR00773	GRPE PROTEIN SIGNATURE	9.112e-09 184-211
11	1.V00//2	OVER LYOTETH SIGNATORE	PR00773D 16.14 5.922e-09 215-235

Table 3

SEQ ID NO:	Database entry ID	Description	*Results
23	PD00930	PROTEIN GTPASE DOMAIN	PD00930B 33.72 7.300e-26 600-203
23	1000930	ACTIVATION.	PD00930A 25.62 1.514e-16 497-523
23	BL50002	Src homology 3 (SH3) domain proteins profile.	BL50002A 14.19 4.000e-12 727-746
23	PF00182	GTPase-activator protein for Rho-like GTPases	PF00182B 14.20 7.333e-12 549-128
25	BL00375	UDP-glycosyltransferases proteins.	BL00375F 16.99 7.061e-35 291-336
			BL00375C 18.27 2.615e-19 126-150
		· ·	BL00375D 14.56 9.000e-17 192-220
	ľ	•	BL00375B 21.22 8.627e-16 67-108
			BL00375G 13.01 4.577e-13 390-430
28	BL01170	Ribosomal protein L6e proteins.	BL01170A 12.34 9.143e-40 139-175
28	PD01457	RIBOSOMAL PROTEIN 40S ZINC- FINGER METAL.	PD01457A 16.51 9.845e-09 67-112
29	BL00359	Ribosomal protein L11 proteins.	BL00359B 23.07 4.231e-24 56-97
		The second secon	BL00359C 22.18 6.148e-22 111-145
		·	BL00359A 20.66 4.000e-21 20-56
29	BL01108	Ribosomal protein L24 proteins.	BL01108A 20.33 1.000e-08 40-73
30	PR00983	CYSTEINYL-TRNA SYNTHETASE	PR00983D 14.16 3.209e-23 270-292
		SIGNATURE	PR00983C 11.27 3.415e-21 239-258
	٠ .		PR00983A 11.10 1.878e-12 75-87
30	BL00178	Aminoacyl-transfer RNA synthetases	BL00178B 7.11 2.286e-09 314-325
		class-I proteins.	
31	PR00718	PHOSPHOLIPASE D SIGNATURE	PR00718E 8.61 1.000e-08 327-351
32	BL00518	Zinc finger, C3HC4 type (RING finger), proteins.	BL00518 12.23 6.133e-10 49-58
33	PF00992	Troponin.	PF00992A 16.67 7.972e-10 10-45 PF00992A 16.67 5.145e-09 17-52 PF00992A 16.67 6.684e-09 56-91
34	BL01019	ADP-ribosylation factors family proteins.	BL01019A 13.20 8.000e-11 68-108
34	PR00449	TRANSFORMING PROTEIN P21	PR00449C 17.27 4.938e-20 75-98
		RAS SIGNATURE	PR00449A 13.20 1.900e-15 34-56
	i		PR00449E 13.50 6.870e-15 173-196
	ļ		PR00449B 14.34 1.360e-10 57-74
			PR00449D 10.79 5.364e-09 137-151
37	PR00764	COMPLEMENT C9 SIGNATURE	PR00764F 16.89 7.783e-11 204-225
37	DM01077	SEX HORMONE-BINDING GLOBULIN.	DM01077A 16.30 1.165e-10 43-90
37	BL00279	Membrane attack complex components / perforin proteins.	BL00279E 37.11 9.163e-09 187-235
38	PR00832	PAXILLIN SIGNATURE	PR00832B 9.87 6.284e-10 768-792
38	PR00806	VINCULIN SIGNATURE	PR00806A 6.63 9.260e-09 766-777
38	PR00049	WILM'S TUMOUR PROTEIN	PR00049D 0.00 8.661e-15 766-781
. =		SIGNATURE	PR00049D 0.00 3.250e-12 764-779
	}		PR00049D 0.00 7.277e-11 765-780
	1		PR00049D 0.00 8.786e-10 763-778
	1		PR00049D 0.00 9.390e-09 762-777
40	BL00226	Intermediate filaments Proteins.	BL00226D 19.10 3.172e-34 397-444
			BL00226B 23.86 5.929e-23 230-278

Table 3

SEQ ID NO:	Database entry ID	Description	*Results
NO.	chuy no		BL00226C 13.23 4.808e-21 296-327
	ļ		BL00226A 12.77 5.065e-13 129-144
	1		BL00226B 23.86 6.400e-10 181-229
41	BL00243	Integrins beta chain cysteine-rich	BL00243I 31.77 2.014e-09 156-199
-11	BECOE	domain proteins.	BL00243I 31.77 5.437e-09 159-202
	1	Commit protons.	BL00243I 31.77 5.690e-09 30-73
41	BL01208	VWFC domain proteins.	BL01208B 15.83 5.865e-09 184-199
41	BL00203	Vertebrate metallothioneins proteins.	BL00203 13.94 3.670e-11 66-112 BL00203
71	BEOUZOS	vertebrate incumonations proteins.	13.94 4.659e-11 40-86 BL00203 13.94
			7.429e-11 70-116 BL00203 13.94 9.505e-11
			140-186 BL00203 13.94 2.723e-10 21-67
		•	BL00203 13.94 2.723e-10 61-107 BL00203
			13.94 3.147e-10 105-151 BL00203 13.94
			4.064e-10 22-68 BL00203 13.94 5.213e-10
			161-207 BL00203 13.94 6.457e-10 26-72
	1		BL00203 13.94 7.032e-10 184-230 BL00203
			13.94 7.223e-10 80-126 BL00203 13.94
			9.043e-10 130-176 BL00203 13.94 1.735e-
	1	,	09 175-221 BL00203 13.94 3.020e-09 150-
	i		196 BL00203 13.94 3.204e-09 65-111
		,	BL00203 13.94 3.296e-09 95-141 BL00203
			13.94 3.663e-09 135-181 BL00203 13.94
• •			5.041e-09 47-93 BL00203 13.94 5.041e-09
		•	85-131 BL00203 13.94 5.500e-09 100-146
	1		BL00203 13.94 5.867e-09 126-172 BL00203
			13.94 5.959e-09 90-136 BL00203 13.94
			6.694e-09 170-216 BL00203 13.94 6.878e-
	j		09 151-197 BL00203 13.94 6.969e-09 17-63
			BL00203 13.94 7.337e-09 115-161 BL00203
			13.94 7.429e-09 71-117 BL00203 13.94
	1		7.704e-09 171-217 BL00203 13.94 8.531e-
			09 155-201 BL00203 13.94 8.714e-09 165-
			211 BL00203 13.94 9.265e-09 116-162
41	BL00269	Mammalian defensins proteins.	BL00269C 16.52 9.289e-09 28-57
14	BBOOLOS	Transminar dolonoms proteins.	BL00269C 16.52 9.289e-09 72-101
41	PD02283	PROTEIN SPORULATION REPEAT	PD02283C 17.54 5.050e-09 138-166
71	1 202203	PRECU.	PD02283C 17.54 5.175e-09 24-52
	l .	11660.	PD02283C 17.54 5.175e-09 68-96
	1	`	PD02283C 17.54 6.738e-09 113-141
	1	. ,	PD02283C 17.54 7.188e-09 163-191
	J		PD02283C 17.54 7.750e-09 173-201
	1		PD02283C 17.54 7.7306-09 173-201 PD02283C 17.54 7.975e-09 128-156
	1		PD02283C 17.54 8.650e-09 148-176
		·	PD02283C 17.54 9.325e-09 118-146
41	BL00799	Granulins proteins.	BL00799D 12.41 7.661e-09 49-96
-T	DE00/33	Grandina process.	BL00799D 12.41 7.001e-09 49-90 BL00799G 9.41 1.000e-08 39-80
43	BL00291	Prion protein.	BL00799G 9.41 1.000e-06 39-80 BL00291A 4.49 4.414e-09 47-82
44	PF00084	Sushi domain proteins (SCR repeat	PF00084B 9.45 7.188e-10 1549-1561
	F100004	proteins.	LL00004D 2'42 \'\'1006-10 \'\742-1201
44	BL00142	Neutral zinc metallopeptidases, zinc-	BL00142 8.38 2.286e-09 730-741
77	DLUUIAZ	1 1100 da zano metanopepudases, zinc-	DLW0147 0.20 7.7000-02 120-141

Table 3

SEQ ID NO:	Database entry ID	Description	*Results
		binding region proteins.	
44	PR00480	ASTACIN FAMILY SIGNATURE	PR00480B 15.41 3.314e-09 725-744
45	BL00414	Profilin proteins.	BL00414D 15.59 9.182e-10 81-108
48	PR00837	ALLERGEN V5/TPX-1 FAMILY	PR00837D 11.12 6.023e-09 22-36
	į	SIGNATURE	
48	BL01009	Extracellular proteins SCP/Tpx- 1/Ag5/PR-1/Sc7 proteins.	BL01009E 13.50 8.204e-09 21-37
49	BL00284	Serpins proteins.	BL00284A 15.64 2.350e-20 85-109
			BL00284D 16.34 4.240e-19 323-350
	ľ	·	BL00284C 28.56 5.600e-17 216-258
		· ·	BL00284E 19.15 7.500e-14 408-433
	1		BL00284B 17.99 9.379e-13 189-210
50	BL01283	T-box domain proteins.	BL01283A 24.15 2.125e-39 148-196
		- Com Commission Processing	BL01283B 23.17 9.438e-34 208-250
			BL01283D 11.70 7.868e-31 298-331
		*	BL01283C 13.05 8.448e-16 260-274
50	PR00937	T-BOX DOMAIN SIGNATURE	PR00937A 15.25 9.182e-26 156-181
٠.			PR00937D 13.41 7.375e-17 259-274
	1		PR00937B 14.58 8.615e-15 223-237
	1		PR00937E 11.86 8.541e-14 301-315
		•	PR00937F 12.53 1.450e-12 322-331
			PR00937C 10.51 1.000e-11 240-250
50	PR00938	BRACHYURY PROTEIN FAMILY SIGNATURE	PR00938C 8.28 6.547e-09 264-282
50	PR00427	INTERLEUKIN-8 RECEPTOR SIGNATURE	PR00427A 16.30 6.776e-09 416-431
51	PD01270	RECEPTOR FC	PD01270D 24.66 8.054e-09 50-86
50	DT 00027	IMMUNOGLOBULIN AFFIN.	DI 000274 27 CO 2 E42 - 12 101 221
52 52	BL00237	G-protein coupled receptors proteins.	BL00237A 27.68 2.543e-13 181-221
52 ,	PR00245	OLFACTORY RECEPTOR	PR00245A 18.03 7.682e-11 150-172
	DD 00005	SIGNATURE	PR00245C 7.84 5.286e-10 290-306
52	PR00237	RHODOPSIN-LIKE GPCR	PR00237C 15.69 3.700e-09 195-218
	DD 00050	SUPERFAMILY SIGNATURE	PR00237G 19.63 8.535e-09 326-353
53	PR00050	COLD SHOCK PROTEIN	PR00050A 11.28 3.143e-12 42-58
	DI 00050	SIGNATURE	PR00050C 9.82 9.151e-11 85-104
53	BL00352	'Cold-shock' DNA-binding domain	BL00352B 23.66 2.881e-13 71-110
		proteins.	BL00352A 12.19 1.327e-10 42-57
56	BL01173	Lipolytic enzymes G-D-X-G family,	BL01173B 13.27 4.462e-17 140-167
		histidine.	BL01173C 8.98 4.349e-14 182-196
			BL01173A 9.41 1.818e-13 454-467
	ľ.	·	BL01173C 8.98 6.553e-13 495-509
			BL01173A 9.41 8.364e-13 107-120
57	PR00321	GAMMA G-PROTEIN (TRANSDUCIN) SIGNATURE	PR00321C 15.39 2.473e-12 123-141
58	PR00937	T-BOX DOMAIN SIGNATURE	PR00937A 15.25 1.000e-24 117-142
	1		PR00937D 13.41 5.500e-18 220-235
			PR00937B 14.58 5.235e-13 184-198
			PR00937F 12.53 1.450e-12 293-302
		•	PR00937E 11.86 1.918e-12 259-273
			PR00937C 10.51 3.133e-11 201-211

Table 3

SEQ ID NO:	Database entry ID	Description	*Results
58	BL01283	T-box domain proteins.	BL01283A 24.15 1.000e-40 109-157
50	2201203	Took domain protons.	BL01283B 23.17 9.156e-34 169-211
]		BL01283C 13.05 8.286e-17 221-235
	l		BL01283D 11.70 5.709e-11 269-302
58	PR00938	BRACHYURY PROTEIN FAMILY SIGNATURE	PR00938C 8.28 7.384e-09 225-243
59	PD02059	CORE POLYPROTEIN PROTEIN GAG CONTAINS: P.	PD02059A 28.10 2.694e-09 116-157
63	BL00196	Ribosomal protein L30 proteins.	BL00196 34.38 3.250e-15 46-97
64	BL00226	Intermediate filaments proteins.	BL00226B 23.86 1.205e-31 264-312
64	BL01305	moaA / nifB / pqqE family proteins.	BL01305B 10.95 8.875e-09 78-88
68	DM00892	3 RETROVIRAL PROTEINASE.	DM00892C 23.55 6.727e-13 33-67
69	PR00874	FUNGI-IV METALLOTHIONEIN SIGNATURE	PR00874C 4.37 7.214e-10 68-83
69	PD00866	GLYCOPROTEIN PROTEIN SPIKE E2 PRECURSOR PEPLOMER.	PD00866L 3.73 6.564e-10 1-11 PD00866L 3.73 1.443e-09 26-36
69	BL00026	Chitin recognition or binding domain	BL00026 12,95 3.013e-09 48-69
		proteins.	
69	DM01724	kw ALLERGEN POLLEN CIM1 HOL- LI.	DM01724 8.14 3.250e-09 10-30
69	BL01208	VWFC domain proteins.	BL01208B 15.83 6.838e-09 111-126
69	BL00243	Integrins beta chain cysteine-rich	BL00243I 31.77 4.838e-10 106-149
		domain proteins.	BL00243I 31.77 7.221e-10 18-61 BL00243I
		_	31.77 1.761e-09 41-84 BL00243I 31.77
			3.408e-09 31-74 BL00243I 31.77 7.465e-09
			71-114
69	BL00203	Vertebrate metallothioneins proteins.	BL00203 13.94 4.107e-13 66-112 BL00203
			13.94 2.138e-12 92-138 BL00203 13.94
			1.099e-11 28-74 BL00203 13.94 3.176e-11
	:	4	82-128 BL00203 13.94 3.374e-11 87-133
			BL00203 13.94 5.846e-11 77-123 BL00203
			13.94 7.231e-11 102-148 BL00203 13.94
			1.670e-10 97-143 BL00203 13.94 2.532e-10
		_	103-149 BL00203 13.94 5.021e-10 88-134
			BL00203 13.94 7.128e-10 38-84 BL00203
		<u>.</u>	13.94 7.168e-10 107-153 BL00203 13.94
			7.702e-10 73-119 BL00203 13.94 9.426e-10
. ,		•	25-71 BL00203 13.94 1.918e-09 101-147
		·	BL00203 13.94 2.745e-09 27-73 BL00203
·			13.94 4.031e-09 71-117 BL00203 13.94
			4.857e-09 36-82 BL00203 13.94 5.041e-09
			98-144 BL00203 13.94 5.154e-09 6-52
11.			BL00203 13.94 6.418e-09 76-122 BL00203
			13.94 7.980e-09 91-137 BL00203 13.94
		·	8.255e-09 13-59 BL00203 13.94 8.898e-09
			48-94
69	PR00876	NEMATODE METALLOTHIONEIN SIGNATURE	PR00876B 7.66 9.514e-09 80-94
73	PR00875	MOLLUSC METALLOTHIONEIN SIGNATURE	PR00875A 5.83 9.679e-10 17-29

Table 3

SEQ ID NO:	Database entry ID	Description	*Results
74	PR00185	HISTONE H4 SIGNATURE	PR00185B 13.68 8.888e-09 364-384
86	PD00066	PROTEIN ZINC-FINGER METAL- BINDI.	PD00066 13.92 7.000e-13 200-213
86	BL00028	Zinc finger, C2H2 type, domain proteins.	BL00028 16.07 6.850e-13 850-867 BL00028 16.07 1.900e-10 184-201 BL00028 16.07 6.100e-10 371-388 BL00028 16.07 6.914e-09 317-334
86	PR00048	C2H2-TYPE ZINC FINGER SIGNATURE	PR00048B 6.02 7.158e-09 197-207
87	PD02870	RECEPTOR INTERLEUKIN-1 PRECURSOR.	PD02870D 15.74 8.468e-09 358-393
88	BL00048	Protamine P1 proteins.	82 BL00048 6.39 5.500e-10 70-97 BL00048 6.39 2.350e-09 62-89 BL00048 6.39 3.700e-09 60-87 BL00048 6.39 5.050e-09 63-90 BL00048 6.39 6.288e-09 61-88 BL00048 6.39 9.438e-09 71-98
89	PR00320	G-PROTEIN BETA WD-40 REPEAT SIGNATURE	PR00320C 13.01 8.920e-10 202-217 PR00320B 12.19 9.486e-10 202-217 PR00320C 13.01 7.900e-09 292-307 PR00320A 16.74 8.902e-09 202-217
90	BL00453	FKBP-type peptidyl-prolyl cis-trans isomerase proteins.	BL00453B 23.86 3.864e-28 106-140 BL00453A 15.57 1.000e-15 81-96 BL00453C 9.72 1.000e-12 147-160
92	PR00299	ALPHA CRYSTALLIN SIGNATURE	PR00299B 17.53 7.211e-09 324-337
93	PF00676	Dehydrogenase E1 component.	PF00676D 14.40 4.857e-13 421-441 PF00676C 16.88 1.931e-10 389-413 PF00676B 24.71 5.433e-10 192-230
96	BL00824	Elongation factor 1 beta/beta'/delta chain proteins.	BL00824B 9.21 3.919e-09 1472-1492
99	PR00417	PROKARYOTIC DNA TOPOISOMERASE I SIGNATURE	PR00417A 12.66 5.415e-09 866-880
102	PD01066	PROTEIN ZINC FINGER ZINC- FINGER METAL-BINDING NU.	PD01066 19.43 6.936e-29 17-56
102	BL00028	Zinc finger, C2H2 type, domain proteins.	BL00028 16.07 2.059e-14 435-452 BL00028 16.07 7.353e-14 351-368 BL00028 16.07 2.350e-13 295-312 BL00028 16.07 9.100e-13 491-508 BL00028 16.07 2.174e-12 463-480 BL00028 16.07 8.826e-12 211-228
			BL00028 16.07 2.038e-11 379-396 BL00028 16.07 2.385e-11 323-340 BL00028 16.07 3.423e-11 239-256 BL00028 16.07 9.654e-11 407-424 BL00028 16.07 1.000e-10 267-284
102	BL00479	Phorbol esters / diacylglycerol binding domain proteins.	BL00479A 19.86 6.362e-09 366-389
102	PD02462	PROTEIN BOLA TRANSCRIPTION REGULATION AC.	PD02462A 22.48 7.695e-09 204-239
102	PR00048	C2H2-TYPE ZINC FINGER SIGNATURE	PR00048A 10.52 1.000e-15 460-474 PR00048A 10.52 1.000e-14 432-446 PR00048A 10.52 3.250e-14 320-334

Table 3

SEQ ID NO:	Database entry ID	Description	*Results
	1 02.02		PR00048A 10.52 4.750e-14 348-362
	•	<u>.</u> .	PR00048A 10.52 6.250e-14 376-390
			PR00048A 10.52 3.133e-13 292-306
	ļ :		PR00048A 10.52 1.529e-12 488-502
			PR00048B 6.02 1.000e-11 336-346
]	:	PR00048B 6.02 9.308e-11 224-234
	1		PR00048B 6.02 2.688e-10 476-486
	!		PR00048B 6.02 3.250e-10 280-290
•			PR00048A 10.52 5.696e-10 404-418
]		PR00048A 10.52 6.087e-10 264-278
	. !		PR00048B 6.02 6.187e-10 420-430
-			PR00048A 10.52 7.214e-10 236-250
			PR00048B 6.02 8.875e-10 364-374
	[,		PR00048B 6.02 3.368e-09 171-181
		•	PR00048B 6.02 4.316e-09 448-458
103	PD01066	PROTEIN ZINC FINGER ZINC-	PD01066 19.43 9.438e-37 10-49
100	1 20.000	FINGER METAL-BINDING NU.	1201000 17:43 7:4300-37 10-47
103	BL00028	Zinc finger, C2H2 type, domain	BL00028 16.07 5.500e-13 413-430 BL00028
103	DE00028	proteins.	16.07 1.000e-12 273-290 BL00028 16.07
	[protens.	1.783e-12 357-374 BL00028 16.07 7.577e-
	1		11 301-318 BL00028 16.07 9.308e-11 441-
		,	458 BL00028 16.07 9.308e-11 441-
			BL00028 16.07 1.300e-10 329-346
103	PR00048	C2H2-TYPE ZINC FINGER	PR00048A 10.52 7.000e-14 354-368
103	1100046	SIGNATURE	PR00048A 10.52 2.286e-13 298-312
•		BIGITATORE	PR00048A 10.52 2.2506=13 298-312
			PR00048A 10.52 3.209e-12 410-424
			PR00048B 6.02 5.000e-12 286-296
			PR00048B 6.02 1.000e-11 342-352
			PR00048B 6.02 1.000e-11 370-380
			PR00048B 6.02 1.125e-10 314-324
		· · · · · · · · · · · · · · · · · · ·	PR00048A 10.52 2.565e-10 466-480
			PR00048A 10.52 4.522e-10 438-452
•			PR00048B 6.02 1.474e-09 454-464
			PR00048A 10.52 3.520e-09 326-340
	l ,	•	PR00048B 6.02 4.789e-09 482-492
103	PD00066	PROTEIN ZINC-FINGER METAL-	PD00066 13.92 8.200e-16 289-302 PD00066
105	1200000	BINDI.	13.92 3.769e-15 317-330 PD00066 13.92
		BIADI.	6.538e-15 373-386 PD00066 13.92 2.800e-
	{		14 345-358 PD00066 13.92 4.600e-14 457-
•		·	470 PD00066 13.92 4.130e-11 401-414
	,		PD00066 13.92 9.654e-10 429-442 PD00066
103	BL01024	Protein phosphatase 2A regulatory	13.92 5.200e-09 261-274
		subunit PR55 proteins.	BL01024H 13.88 7.353e-09 163-216
104	PD01781	PROTEASE IMMUNOGLOBULIN PRECURSO.	PD01781B 27.55 8.680e-09 325-369
105	PD01781	PROTEASE IMMUNOGLOBULIN	PD01781B 27.55 8.680e-09 379-423
107	DDOCCOC	PRECURSO.	DD 00000D 10 07 0 000
107	PR00939	C2HC-TYPE ZINC-FINGER	PR00939B 13.27 3.209e-09 1302-1311

Table 3

SEQ ID NO:	Database entry ID	Description	*Results
		SIGNATURE	
108	PD00066	PROTEIN ZINC-FINGER METAL-	PD00066 13.92 2.800e-14 279-292 PD00066
		BINDI.	13.92 4.600e-14 307-320 PD00066 13.92
		· '-	1.000e-13 335-348 PD00066 13.92 7.500e-
	·		13 363-376
108	BL00028	Zinc finger, C2H2 type, domain	BL00028 16.07 7.882e-14 319-336 BL00028
•		proteins.	16.07 7.300e-13 347-364 BL00028 16.07
			4.913e-12 291-308 BL00028 16.07 2.500e-
			10 263-280 BL00028 16.07 1.257e-09 375-
			392
108	PR00048	C2H2-TYPE ZINC FINGER	PR00048A 10.52 4.214e-13 288-302
		SIGNATURE	PR00048B 6.02 5.000e-12 304-314
			PR00048A 10.52 6.824e-12 372-386
		. •	PR00048A 10.52 7.353e-12 344-358
*			PR00048A 10.52 7.158e-11 316-330
			PR00048B 6.02 7.231e-11 276-286
	j		PR00048B 6.02 1.000e-09 332-342
100			PR00048B 6.02 6.211e-09 388-398
108	BL00115	Eukaryotic RNA polymerase II	BL00115Z 3.12 8.842e-18 96-145
		heptapeptide repeat proteins.	BL00115Z 3.12 7.144e-17 89-138
		·	BL00115Z 3.12 6.888e-16 103-152
			BL00115Z 3.12 7.791e-15 82-131
			BL00115Z 3.12 3.947e-14 61-110
			BL00115Z 3.12 7.292e-14 117-166
	•		BL00115Z 3.12 9.164e-14 110-159
			BL00115Z 3.12 1.000e-13 75-124
		·	BL00115Z 3.12 3.871e-13 54-103
		·	BL00115Z 3.12 6.819e-13 68-117
			BL00115Z 3.12 4.168e-11 124-173 BL00115Z 3.12 9.651e-10 47-96 BL00115Z
			3.12 7.485e-09 71-120 BL00115Z 3.12
•		,	9.669e-09 78-127
109	PR00193	MYOSIN HEAVY CHAIN	PR00193D 14.36 5.680e-33 391-420
107	1100175	SIGNATURE	PR00193D 14.30 3.060e-33 391-420
		DIGITIONE	PR00193B 11.69 1.692e-26 110-136
		·	PR00193E 19.47 5.500e-21 445-474
			PR00193A 15.41 4.130e-20 54-74
		•	PR00193E 19.47 5.091e-12 444-473
110	BL00239	Receptor tyrosine kinase class II	BL00239B 25.15 2.985e-16 67-115
		proteins.	DE00237B 23.13 2.7030-10 07-113
110	PR00109	TYROSINE KINASE CATALYTIC	PR00109B 12.27 8.660e-13 132-151
		DOMAIN SIGNATURE	11.001030 12.27 0.0000-13 132-131
110	BL00107	Protein kinases ATP-binding region	BL00107A 18.39 4.462e-25 132-163
		proteins.	BL00107B 13.31 6.143e-10 197-213
110	DM00406	GLIADIN.	DM00406 7.73 1.800e-09 818-831
110	BL00904	Protein prenyltransferases alpha subunit	BL00904A 8.30 5.596e-09 815-865
		repeat proteins proteins.	
110	BL00415	Synapsins proteins.	BL00415A 6.15 7.684e-09 796-837
110	DM00215	PROLINE-RICH PROTEIN 3.	
110	DM00215	PROLINE-RICH PROTEIN 3.	DM00215 19.43 2.373e-09 801-834 DM00215 19.43 7.712e-09 797-830

Table 3

SEQ ID NO:	Database entry ID	Description	*Results
110	PR00209	ALPHA/BETA GLIADIN FAMILY SIGNATURE	PR00209B 4.88 4.188e-09 817-836 PR00209C 4.56 8.929e-09 790-804
111	BL00678	Trp-Asp (WD) repeat proteins proteins.	BL00678 9.67 2.800e-10 366-377 BL00678
İ	Ī	' ' ' ' ' ' ' ' ' ' ' ' ' ' ' ' ' ' '	9.67 5.263e-09 417-428 BL00678 9.67
]			6.211e-09 186-197
111	PR00308	TYPE I ANTIFREEZE PROTEIN	PR00308C 3.83 8.892e-10 108-118
ļ		SIGNATURE	PR00308C 3.83 8.892e-10 109-119
			PR00308C 3.83 8.364e-09 107-117
111	PR00320	G-PROTEIN BETA WD-40 REPEAT	PR00320A 16.74 4.000e-13 364-379
		SIGNATURE	PR00320B 12.19 7.923e-12 415-430
	,		PR00320A 16.74 5.966e-11 415-430
		·	PR00320C 13.01 7.214e-11 415-430
	1		PR00320C 13.01 9.217e-11 364-379
			PR00320A 16.74 9.690e-11 184-199
			PR00320B 12.19 3.057e-10 184-199
	}		PR00320C 13.01 6.040e-10 184-199
•		÷	PR00320B 12.19 6.657e-10 364-379
			PR00320B 12.19 1.450e-09 457-472
•			PR00320C 13.01 2.200e-09 240-255
			PR00320A 16.74 4.732e-09 457-472
			PR00320A 16.74 6.488e-09 281-296
			PR00320C 13.01 1.000e-08 281-296
112	DM00547	1 kw CHROMO BROMODOMAIN	DM00547F 23.43 2.350e-35 384-431
		SHADOW GLOBAL.	DM00547C 17.30 7.000e-19 23-45
٠.			DM00547E 13.94 5.154e-17 135-158
110	DT 00015	7	DM00547D 11.60 2.750e-13 105-119
112	BL00315	Dehydrins proteins.	BL00315A 9.35 4.246e-10 1301-1329
112	PF00426	Outer Capsid protein VP4 (Hemagglutinin).	PF00426S 15.67 6.438e-10 1271-1309
112	BL00039	DEAD-box subfamily ATP-dependent	BL00039D 21.67 6.793e-10 368-414
		helicases proteins.	
112	PD02191	I ATP-BINDING NUCLEOSIDE TRANSCR.	PD02191A 13.95 9.036e-10 107-122
112	BL00048	Protamine P1 proteins.	BL00048 6.39 1.900e-09 1257-1284
		· ·	BL00048 6.39 5.050e-09 1258-1285
112	PF00774	Dihydropyridine sensitive L-type	PF00774A 16.47 7.130e-09 1280-1326
		calcium channel (Beta subuni.	PF00774A 16.47 7.730e-09 1276-1322
112	BL00115	Eukaryotic RNA polymerase II	BL00115Z 3.12 3.448e-11 1254-1303
		heptapeptide repeat proteins.	BL00115Z 3.12 3.302e-10 1261-1310
		· · · · · · · · · · · · · · · · · · ·	BL00115Z 3.12 4.837e-10 1258-1307
	1		BL00115Z 3.12 7.767e-10 1251-1300
			BL00115Z 3.12 8.167e-10 1263-1312
			BL00115Z 3.12 8.884e-10 1260-1309 09
			1247-1296 BL00115Z 3.12 2.985e-09 1240-
		·	1289 BL00115Z 3.12 5.632e-09 1265-1314
	İ		BL00115Z 3.12 8.676e-09 1253-1302
			BL00115Z 3.12 9.471e-09 1268-1317
444			BL00115Z 3.12 9.735e-09 1257-1306
112	PF00186	Flocculin repeat proteins.	PF00186I 9.10 5.290e-13 1279-1309
			PF00186I 9.10 6.838e-12 1277-1307

Table 3

SEQ ID NO:	Database entry ID	Description	*Results
			PF00186I 9.10 2.957e-11 1282-1312
	Ì		PF00186I 9.10 7.496e-11 1276-1306
			PF00186I 9.10 5.200e-10 1268-1298
		· ·	PF00186I 9.10 7.450e-10 1278-1308
			PF00186I 9.10 7.450e-10 1280-1310
ŀ			PF00186I 9.10 4.543e-09 1266-1296
İ			PF00186I 9.10 5.252e-09 1285-1315
ļ		·	PF00186I 9.10 6.031e-09 1272-1302
	1		PF00186I 9.10 6.102e-09 1274-1304
	ļ		PF00186I 9.10 7.236e-09 1270-1300
		·	PF00186I 9.10 8.016e-09 1261-1291
			PF00186I 9.10 9.433e-09 1262-1292
			PF00186I 9.10 9.433e-09 1267-1297
			PF00186I 9.10 1.000e-08 1256-1286
114	PR00700	PROTEIN TYROSINE	PR00700D 12.47 8.788e-11 237-256
		PHOSPHATASE SIGNATURE	
114	BL00383	Tyrosine specific protein phosphatases proteins.	BL00383E 10.35 5.327e-10 240-251
116	PR00884	RIBOSOMAL PROTEIN HS6	PR00884E 8.32 4.750e-09 449-466
		SIGNATURE	
117	PD02890	ISOMERASE CHALCONE-	PD02890C 16.14 8.457e-09 200-235
		FLAVONONE FLAV.	
118	BL00226	Intermediate filaments proteins.	BL00226B 23.86 6.513e-10 401-449
118	BL00326	Tropomyosins proteins.	BL00326D 8.76 1.925e-09 196-237
118	BL01160	Kinesin light chain repeat proteins.	BL01160B 19.54 2.678e-09 328-382
			BL01160B 19.54 8.932e-09 654-708
119	PD01823	PROTEIN INTERGENIC REGION	PD01823C 16.13 7.000e-14 352-373
		ABC1 PRECURSOR	PD01823B 14.96 3.782e-13 328-348
		MITOCHONDRION T.	PD01823D 16.66 6.857e-10 430-451
119	PD01115	PRECURSOR AMPHIBIAN SKIN SIGNAL.	PD01115B 12.92 8.421e-09 268-282
122	BL00854	Proteasome B-type subunits proteins.	BL00854C 29.92 8.435e-19 114-143
124	BL00651	Ribosomal protein L9 proteins.	BL00651A 23.25 8.477e-17 94-134
125	BL01245	RIO1/ZK632.3/MJ0444 family	BL01245F 18.75 2.373e-23 334-371
		proteins.	BL01245A 14.04 8.342e-23 206-231
	[.	BL01245C 13.31 6.564e-15 262-282
			BL01245E 15.28 1.000e-12 320-330
•			BL01245B 11.91 9.809e-10 245-255
128	PR00793	PROLYL AMINOPEPTIDASE (S33) FAMILY SIGNATURE	PR00793C 12.24 1.333e-09 168-183
128	PR00111	ALPHA/BETA HYDROLASE FOLD SIGNATURE	PR00111C 13.46 6.000e-09 182-196
129	BL01160	Kinesin light chain repeat proteins.	BL01160D 10.17 7.077e-09 505-534
129	PD00126	PROTEIN REPEAT DOMAIN TPR NUCLEA.	PD00126A 22.53 1.000e-08 695-716
130	BL00355	HMG14 and HMG17 proteins.	BL00355 5.97 8.412e-32 18-49
130	PR00925	NONHISTONE CHROMOSOMAL	PR00925B 3.73 3.400e-16 34-47 PR00925A
		PROTEIN HMG17 FAMILY	5.47 1.750e-15 18-33 PR00925C 5.57
		SIGNATURE	9.824e-09 51-62
131	PR00041	CAMP RESPONSE ELEMENT	PR00041E 7.20 2.976e-13 305-326

Table 3

SEQ ID NO:	Database entry ID	Description	*Results
		BINDING (CREB) PROTEIN SIGNATURE	
131	BL00036	bZIP transcription factors basic domain proteins.	BL00036 9.02 4.103e-09 299-312
132	PR00211	GLUTELIN SIGNATURE	PR00211B 0.86 1.750e-09 205-226
			PR00211B 0.86 8.750e-09 199-220
132	DM00215	PROLINE-RICH PROTEIN 3.	DM00215 19.43 4.529e-11 201-234
•			DM00215 19.43 1.804e-10 195-228
		· ·	DM00215 19.43 2.768e-10 192-225
]	•	DM00215 19.43 4.054e-10 202-235
			DM00215 19.43 6.304e-10 207-240
			DM00215 19.43 7.429e-10 180-213
		, .	DM00215 19.43 8.393e-10 196-229
			DM00215 19.43 8.714e-10 218-251
	1		DM00215 19.43 6.034e-09 185-218
		·	DM00215 19.43 6.034e-09 219-252
]		DM00215 19.43 6.492e-09 223-256
	i	'	DM00215 19.43 7.254e-09 200-233
		•	DM00215 19.43 9.390e-09 189-222
122	DY 00455	D / / A / CO 1 : 1 : 1	DM00215 19.43 9.695e-09 213-246
133	BL00455	Putative AMP-binding domain proteins.	BL00455 13.31 5.125e-11 293-309
133	PR00154	AMP-BINDING SIGNATURE	PR00154A 8.88 6.276e-09 286-298
136	PD00015	GLYCOPROTEIN PRECURSOR CELL SI.	PD00015A 8.90 6.400e-09 243-251
138	BL00227	Tubulin subunits alpha, beta, and	BL00227B 19.29 1.000e-40 52-107
·		gamma proteins.	BL00227C 25.48 1.000e-40 113-165
	_		BL00227A 24.55 8.200e-36 1-35
140	PR00049	WILM'S TUMOUR PROTEIN	PR00049D 0.00 8.377e-13 60-75 PR00049D
		SIGNATURE	0.00 7.500e-10 63-78 PR00049D 0.00
			8.071e-10 61-76
140	PR00806	VINCULIN SIGNATURE	PR00806B 4.28 8.440e-09 68-82
140	BL00904	Protein prenyltransferases alpha subunit repeat proteins proteins.	BL00904A 8.30 9.553e-09 60-110
141	PR00049	WILM'S TUMOUR PROTEIN SIGNATURE	PR00049D 0.00 6.438e-12 1175-1190
141	BL01187	Calcium-binding EGF-like domain	BL01187B 12.04 5.800e-11 1284-1300
		proteins pattern proteins.	BL01187B 12.04 8.200e-11 180-196
141	BL01248	Laminin-type EGF-like (LE) domain	BL01248 11.02 4.343e-12 1362-1375
		proteins.	BL01248 11.02 2.350e-11 322-335 BL01248
			11.02 4.125e-10 271-284
141	PR00764	COMPLEMENT C9 SIGNATURE	PR00764B 13.56 3.475e-09 1047-1068
141	PR00010	TYPE II EGF-LIKE SIGNATURE	PR00010C 11.16 4.205e-09 185-196
141	BL01113	C1q domain proteins.	BL01113A 17.99 5.673e-09 1621-1210
141	PR00011	TYPE III EGF-LIKE SIGNATURE	PR00011D 14.03 8.895e-12 551-132
			PR00011B 13.08 5.846e-11 551-132
	· [•	PR00011D 14.03 3.215e-10 313-332
			PR00011A 14.06 4.214e-10 313-332
		. •	PR00011B 13.08 7.783e-10 313-332
			PR00011A 14.06 7.781e-09 551-132
141	BL00420	Speract receptor repeat proteins domain	BL00420A 20.42 8.200e-09 1186-1215

Table 3

SEQ ID NO:	Database entry ID	Description	*Results
		proteins.	
141	PD02510	ISOMERASE GALACTOSE-6- PHOSPHATE.	PD02510B 18.31 8.170e-09 548-144
141	PR00261	LOW DENSITY LIPOPROTEIN (LDL) RECEPTOR SIGNATURE	PR00261F 11.57 9.544e-09 1052-1074
141	PR00288	PUROTHIONIN SIGNATURE	PR00288C 10.15 9.165e-09 311-326
142	DM01970	0 kw ZK632.12 YDR313C ENDOSOMAL III.	DM01970B 8.60 4.750e-17 114-565
142	BL01160	Kinesin light chain repeat proteins.	BL01160B 19.54 2.373e-09 203-257
142	BL00518	Zinc finger, C3HC4 type (RING finger), proteins.	BL00518 12.23 4.000e-09 559-130
142	BL00422	Granins proteins.	BL00422E 26.86 8.615e-09 462-498
143	PD00066	PROTEIN ZINC-FINGER METAL- BINDI.	PD00066 13.92 5.846e-15 141-154 PD00066 13.92 9.217e-11 551-564 PD00066 13.92 6.700e-09 523-536
143	PR00048	C2H2-TYPE ZINC FINGER SIGNATURE	PR00048A 10.52 9.526e-11 122-136 PR00048A 10.52 2.174e-10 532-546 PR00048A 10.52 6.087e-10 588-164 PR00048B 6.02 7.632e-09 138-148 PR00048A 10.52 8.920e-09 504-518
143	PF00651	BTB (also known as BR-C/Ttk) domain proteins.	PF00651 15.00 8.920e-09 59-72
143	BL00028	Zinc finger, C2H2 type, domain proteins.	BL00028 16.07 7.577e-11 535-114 BL00028 16.07 2.200e-10 125-142 BL00028 16.07 5.800e-10 507-524 BL00028 16.07 8.714e-09 591-170 BL00028 16.07 9.743e-09 444-461
144	PR00926	MITOCHONDRIAL CARRIER PROTEIN SIGNATURE	PR00926F 17.75 3.672e-10 262-285
144	BL00215	Mitochondrial energy transfer proteins.	BL00215A 15.82 7.900e-15 16-41 BL00215A 15.82 8.147e-14 260-285 BL00215A 15.82 1.804e-09 166-191 BL00215B 10.44 5.500e-09 114-127
144	PR00927	ADENINE NUCLEOTIDE TRANSLOCATOR 1 SIGNATURE	PR00927B 14.66 8.644e-09 104-126
147	DM01417	6 kw INDUCING XPMC2 MUSHROOM SPAC22G7.04.	DM01417C 12.93 3.250e-11 267-279 DM01417D 11.08 2.200e-10 306-322
148	BL01160	Kinesin light chain repeat proteins.	BL01160B 19.54 8.378e-10 349-403
151	PR00049	WILM'S TUMOUR PROTEIN SIGNATURE	PR00049D 0.00 7.807e-11 419-434 PR00049D 0.00 8.125e-11 1284-1299 PR00049D 0.00 3.929e-10 1283-1298 PR00049D 0.00 3.288e-09 417-432
151	BL00904	Protein prenyltransferases alpha subunit repeat proteins proteins.	BL00904A 8.30 3.553e-09 416-466
154	BL00665	Dihydrodipicolinate synthetase proteins.	BL00665D 14.76 1.000e-11 109-132 BL00665C 25.58 5.832e-11 50-101
154	PR00146	DIHYDRODIPICOLINATE SYNTHASE SIGNATURE	PR00146D 16.26 2.525e-10 108-126 PR00146A 12.62 8.615e-09 13-35
156	PD02906	SYNTHASE I PSEUDOURIDYLATE PSEUDOURIDINE LYASE TR.	PD02906C 24.17 9.115e-15 171-206 PD02906B 15.35 4.886e-13 142-155

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Table 3

SEQ ID NO:	Database entry ID	Description	*Results
			PD02906D 12.27 1.000e-09 239-249
			PD02906A 10.84 8.333e-09 92-105
157	BL00107	Protein kinases ATP-binding region	BL00107B 13.31 2.286e-11 396-412
	ļ	proteins.	BL00107A 18.39 6.148e-11 332-363
157	PR00109	TYROSINE KINASE CATALYTIC DOMAIN SIGNATURE	PR00109B 12.27 4.938e-09 332-351
160	PF01008	Initiation factor 2 subunit.	PF01008B 25.59 9.171e-36 366-409
		•	PF01008A 20.14 8.676e-12 315-336
	<u> </u>		PF01008C 12.25 7.382e-10 449-469
161	BL00591	Glycosyl hydrolases family 10 proteins.	BL00591D 8.33 6.167e-09 2099-2112
163	PR00019	LEUCINE-RICH REPEAT	PR00019B 11.36 7.120e-09 99-113
	<u> </u>	SIGNATURE	PR00019B 11.36 7.840e-09 73-87
164	BL00198	Nt-dnaJ domain proteins.	BL00198A 8.07 3.000e-14 143-160
164	PR00187	DNAJ PROTEIN FAMILY SIGNATURE	PR00187A 12.84 8.800e-12 139-159
165	PR00310	ANTI-PROLIFERATIVE PROTEIN	PR00310B 10.59 4.000e-39 41-71
		BTG1 FAMILY SIGNATURE	PR00310C 12.74 2.256e-33 71-101
	Ī		PR00310D 9.10 9.820e-33 101-131
			PR00310A 11.17 7.000e-27 16-41
165	BL00960	BTG1 family proteins.	BL00960B 24.47 1.000e-40 34-79
l		•	BL00960C 12.68 6.745e-21 98-120
			BL00960A 10.98 5.304e-12 14-26
166	BL00216	Sugar transport proteins.	BL00216B 27.64 2.688e-21 124-174
166	DM00973	3 kw RESISTANCE BENOMYL YLL028W CYCLOHEXIMIDE.	DM00973A 21.17 4.162e-10 96-133
166	PR00171	SUGAR TRANSPORTER	PR00171D 12.76 3.520e-13 456-478
100	PROOL/I	SIGNATURE	PR00171D 12.76 3.3206-13 436-478 PR00171E 14.87 2.750e-09 479-492
166	PR00172	GLUCOSE TRANSPORTER	PR00172D 9.13 6.513e-09 456-480
100	1100172	SIGNATURE	BL00216B 27.64 5.198e-20 124-174
167	BL00216	Sugar transport proteins.	
167	DM00973	3 kw RESISTANCE BENOMYL	DM00973A 21.17 4.162e-10 96-133
		YLL028W CYCLOHEXIMIDE.	
168	PD01066	PROTEIN ZINC FINGER ZINC-	PD01066 19.43 5.929e-32 59-98
		FINGER METAL-BINDING NU.	·
168	PD00066	PROTEIN ZINC-FINGER METAL-	PD00066 13.92 2.385e-15 520-533 PD00066
	1	BINDL	13.92 2.800e-14 296-309 PD00066 13.92
			5.200e-14 240-253 PD00066 13.92 5.200e-
	1		14 548-561 PD00066 13.92 9.400e-14 436-
•] :		449 PD00066 13.92 1.000e-13 324-337
		•	PD00066 13.92 6.143e-12 352-365 PD00066
			13.92 6.885e-10 268-281
168	PR00048	C2H2-TYPE ZINC FINGER	PR00048B 6.02 6.000e-12 237-247
		SIGNATURE	PR00048A 10.52 6.294e-12 333-347
			PR00048A 10.52 6.824e-12 361-375
	1		PR00048A 10.52 9.471e-12 249-263
	1		PR00048A 10.52 4.316e-11 119-133
	1		PR00048A 10.52 4.789e-11 529-543
	1		PR00048A 10.52 6.684e-11 445-459
	1		PR00048A 10.52 8.141e-11 305-319
		<u> </u>	PR00048B 6.02 6.063e-10 321-331

Table 3

SEQ ID	Database	Description	*Results
NO:	entry ID		DD00040D C00 C002 10 517 507
]	}		PR00048B 6.02 6.063e-10 517-527
			PR00048A 10.52 7.261e-10 221-235
1	1	·	PR00048B 6.02 7.750e-10 545-117
[PR00048B 6.02 1.474e-09 293-303
			PR00048A 10.52 2.800e-09 389-403
	7700455		PR00048A 10.52 1.000e-08 417-431
170	PR00456	RIBOSOMAL PROTEIN P2	PR00456E 3.06 2.820e-11 6-21 PR00456E
	770000	SIGNATURE	3.06 7.125e-10 3-18
170	PD02331	CYCLIN CELL CYCLE DIVISION	PD02331A 19.76 7.429e-15 93-140
		PROTE.	PD02331B 13.43 1.125e-09 174-207
170	PR00833	POLLEN ALLERGEN POA PI	PR00833H 2.30 5.269e-09 3-18
		SIGNATURE	
171	PD00126	PROTEIN REPEAT DOMAIN TPR	PD00126A 22.53 4.706e-14 140-161
	<u> </u>	NUCLEA.	PD00126A 22.53 6.824e-14 289-310
173	BL00741	Guanine-nucleotide dissociation	BL00741B 14.27 3.418e-11 294-317
	<u></u>	stimulators CDC24 family sign.	
173	PR00452	SH3 DOMAIN SIGNATURE	PR00452B 11.65 5.154e-11 86-102
173	PR00497	NEUTROPHIL CYTOSOL FACTOR	PR00497D 11.91 5.962e-10 91-113
		P40 SIGNATURE	
173	PF00564	Octicosapeptide repeat proteins.	PF00564B 24.74 6.442e-09 277-328
175	BL01016	Glycoprotease family proteins.	BL01016C 22.84 5.292e-19 60-105
	,		BL01016H 13.71 6.157e-12 307-317
			BL01016E 14.88 3.182e-11 141-169
			BL01016D 8.86 6.741e-09 118-131
175	PR00789	O-SIALOGLYCOPROTEIN	PR00789E 12.42 7.128e-14 141-163
·	•	ENDOPEPTIDASE (M22)	PR00789C 16.11 2.707e-12 85-105
		METALLO-PROTEASE FAMILY	PR00789B 10.48 1.205e-09 64-85
		SIGNATURE	PR00789D 8.17 7.151e-09 118-131
176	PR00850	GLYCOSYL HYDROLASE FAMILY	PR00850B 6.67 5.455e-09 148-173
		59 SIGNATURE	
178	PR00259	TRANSMEMBRANE FOUR FAMILY	PR00259A 9.27 8.676e-20 17-41 PR00259C
		SIGNATURE	16.40 4.750e-17 85-114 PR00259B 14.81
			8.615e-12 58-85 PR00259D 13.50 2.528e-11
			235-262
178	BL00421	Transmembrane 4 family proteins.	BL00421B 17.62 6.186e-17 64-103
		Transmission 4 minsy process.	BL00421A 11.79 6.800e-12 13-32
			BL00421E 20.97 1.514e-10 232-262
			BL00421C 12.89 3.600e-09 147-159
178	PR00235	HERPESVIRUS MAJOR CAPSID	PR00235A 14.64 8.000e-09 87-111
	1100255	PROTEIN (MCP) SIGNATURE	11K00255A 14.04 6.0006-05 67-111
179	BL01052	Calponin family repeat proteins.	BL01052C 18.51 6.806e-40 87-127
	220102	Carpoint tainty topeat protests.	BL01052A 16.12 7.180e-32 3-35 BL01052B
		•	15.31 8.031e-26 52-78 BL01052D 10.26
			1.000e-24 174-194
179	PR00890	SMOOTH MUSCLE PROTEIN 22-	PR00890E 14.34 3.813e-21 135-155
-17	1100000	ALPHA (TRANSGELIN)	PR00890E 14.54 5.815e-21 155-155 PR00890A 8.61 9.775e-21 34-54 PR00890C
		SIGNATURE	8.22 1.000e-17 84-98 PR00890B 8.75
		DIGITATURE	3.455e-17 62-78 PR00890F 12.92 4.064e-14
			161-174 PR00890D 16.17 5.174e-13 118-
			128

Table 3

SEQ ID	Database	Description	*Results
NO:	entry ID		
179	PR00888	SMOOTH MUSCLE	PR00888H 9.97 5.154e-20 175-191
		PROTEIN/CALPONIN FAMILY	PR00888C 12.27 5.179e-18 52-68
	}	SIGNATURE	PR00888D 16.09 4.273e-17 88-105
	1.		PR00888A 11.87 2.350e-16 3-18 PR00888E
	Í	<u> </u>	11.81 3.432e-16 104-120 PR00888F 7.44
	İ		4.825e-14 125-140 PR00888G 12.73 8.759e-
			14 162-176 PR00888B 13.72 2.350e-12 22-
	ļ		36
179	PR00889	CALPONIN SIGNATURE	PR00889E 12.18 2.726e-12 171-187
180	BL00875	Bacterial type II secretion system	BL00875A 25.57 6.447e-09 367-399
		protein D proteins.	22000731123.37 0.4470-07 307-377
181	PD01351	PROTEIN REPEAT	PD01351B 13.72 5.355e-09 238-264
		NEUROFILAMENT TRIPL.	12013312 13.72 3.3330-07 230-204
182	DM01354	kw TRANSCRIPTASE REVERSE II	DM01354H 18.00 8.826e-27 109-149
		ORF2.	DM01354G 11.57 2.143e-25 78-109
			DM01354F 14.56 1.414e-15 42-78
	ł		DM01354E 18.69 8.650e-14 17-47
182	BL00869	Renal dipeptidase proteins.	BL00869D 14.02 3.477e-09 67-96
185	BL00039	DEAD-box subfamily ATP-dependent	BL00039A 18.44 4.000e-25 222-261
	2200035	helicases proteins.	BL00039A 18.44 4.000e-23 222-261 BL00039D 21.67 4.529e-23 498-544
		nonouses proteins.	BL00039D 21:07 4:3296-23 496-344 BL00039C 15:63 4:300e-16 347-371
			BL00039C 13.03 4.300e-10 347-371 BL00039B 19.19 9.379e-15 262-288
185	PD00302	PROTEASE POLYPROTEIN	PD00302B 9.52 1.346e-09 234-250
100	1 200302	HYDROLASE ASP.	FD00302B 9.32 1.340e-09 234-230
186	PD00066	PROTEIN ZINC-FINGER METAL-	PD00066 13.92 5.714e-12 152-165 PD00066
	120000	BINDI.	13.92 6.143e-12 124-137
186	BL00028	Zinc finger, C2H2 type, domain	BL00028 16.07 6.885e-11 136-153 BL00028
	2200020	proteins.	16.07 2.200e-10 197-214
186	PR00239	MOLLUSCAN RHODOPSIN C-	PR00239E 1.58 5.705e-09 420-432
		TERMINAL TAIL SIGNATURE	110023711.36 3.7036-03 420-432
186	PR00048	C2H2-TYPE ZINC FINGER	PR00048A 10.52 2.957e-10 133-147
	11000-10	SIGNATURE	PR00048A 10.52 2,937e-10 133-147
		SIGNATURE	PR00048A 10.52 8.043e-10 161-175
;			PR00048B 6.02 8.105e-09 121-131
187	BL01022	PTR2 family proton/oligopeptide	BL01022B 22.19 4.240e-10 308-354
107	2201022	symporters proteins.	BL01022B 22.19 4.240e-10 506-554
187	PR00669	INHIBIN ALPHA CHAIN	PR00669B 8.27 7.915e-09 264-281
10.	1100000	SIGNATURE	FR00009B 6.27 7.9156-09 204-281
190	PR00830	ENDOPEPTIDASE LA (LON)	PR00830A 8.41 3.342e-09 881-901
.,,	1100050	SERINE PROTEASE (S16)	FR00050A 8.41 5.342e-09 881-901
	·	SIGNATURE	
191	PR00109	TYROSINE KINASE CATALYTIC	PRO0100P 12 27 0 224- 12 261 200
-/-	* 1700103	DOMAIN SIGNATURE	PR00109B 12.27 9.234e-13 261-280
191	BL00107		DT 001074 19 20 1 000 - 02 261 000
•/•	יסומתים /	Protein kinases ATP-binding region proteins.	BL00107A 18.39 1.000e-23 261-292
191	BL00239	Receptor tyrosine kinase class II	BL00107B 13.31 1.000e-12 341-357
171	ו אכשונים		BL00239B 25.15 6.523e-10 196-244
191	DI 00470	proteins.	DV 004700 10 01 1 000 00 000
171	BL00479	Phorbol esters / diacylglycerol binding	BL00479C 12.01 1.000e-09 320-333
101	DD00024	domain proteins	PRO00247-10-01-0-045-00-00-00-00-00-00-00-00-00-00-00-00-00
191	PR00834	HTRA/DEGQ PROTEASE FAMILY	PR00834F 10.91 2.946e-09 786-799

Table 3

SEQ ID NO:	Database entry ID	Description	*Results
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193	BL01033	Globins profile.	BL01033A 16.94 2.385e-18 25-47
193	PR00814	BETA HAEMOGLOBIN	PR00814A 12.94 1.000e-22 30-47
		SIGNATURE	PR00814B 9.18 7.750e-18 48-64
193	PR00175	MYOGLOBIN SIGNATURE	PR00175B 9.02 9.392e-10 25-49
194	PR00320	G-PROTEIN BETA WD-40 REPEAT	PR00320B 12.19 6.226e-11 140-155
		SIGNATURE	PR00320A 16.74 4.971e-10 140-155
	L		PR00320C 13.01 9.280e-10 140-155
194	BL00678	Trp-Asp (WD) repeat proteins proteins.	BL00678 9.67 7.632e-09 142-153
196	PR00832	PAXILLIN SIGNATURE	PR00832B 9.87 9.174e-10 309-333
196	BL01160	Kinesin light chain repeat proteins.	BL01160B 19.54 2.054e-10 376-430
			BL01160B 19.54 6.919e-10 383-437
			BL01160B 19.54 9.676e-10 369-423
196	PR00049	WILM'S TUMOUR PROTEIN SIGNATURE	PR00049D 0.00 8.780e-09 40-55
196	BL00087	Copper/Zinc superoxide dismutase proteins.	BL00087C 20.18 8.784e-09 260-296
196	PR00806	VINCULIN SIGNATURE	PR00806A 6.63 9.014e-09 308-319
196	BL00326	Tropomyosins proteins.	BL00326A 14.01 9.143e-09 506-540
197	PR00674	LIGHT HARVESTING PROTEIN B CHAIN SIGNATURE	PR00674A 20.10 7.391e-09 134-155
198	PR00192	F-ACTIN CAPPING PROTEIN BETA	PR00192C 6.65 2.500e-36 57-84 PR00192D
		SUBUNIT SIGNATURE	8.23 4.462e-36 97-125 PR00192E 8.85
			7.000e-33 212-239 PR00192A 8.23 1.474e-
			27 5-26 PR00192B 6.20 3.000e-26 26-48
198	BL00231	F-actin capping protein beta subunit	BL00231A 8.59 1.000e-40 5-51 BL00231B
		proteins.	14.16 1.000e-40 84-128 BL00231D 15.40
		. •	1.000e-40 165-200 BL00231E 11.66 1.000e-
			40 209-246 BL00231C 12.77 1.180e-15 146-
			157
199	PF00023	Ank repeat proteins.	PF00023A 16.03 4.750e-10 45-61
199	PF00791	Domain present in ZO-1 and Unc5-like	PF00791B 28.49 8.768e-12 87-142
		netrin receptors.	PF00791B 28.49 7.028e-09 499-116
199	BL01160	Kinesin light chain repeat proteins.	BL01160E 8.74 7.398e-09 323-362
201	PR00239	MOLLUSCAN RHODOPSIN C-	PR00239E 1.58 6.114e-09 183-195
	·	TERMINAL TAIL SIGNATURE	
202	BL00412	Neuromodulin (GAP-43) proteins.	BL00412D 16.54 4.033e-10 319-370
202	BL00224	Clathrin light chain proteins.	BL00224B 16.94 4.845e-09 313-366
202	PF00992	Troponin.	PF00992A 16.67 8.734e-12 333-368
	1		PF00992A 16.67 2.776e-09 344-379
			PF00992A 16.67 5.026e-09 351-386
203	BL00790	Receptor tyrosine kinase class V proteins.	BL00790R 16.20 7.677e-09 29-73
204	BL00790	Receptor tyrosine kinase class V proteins.	BL00790R 16.20 7.677e-09 29-73
205	BL00790	Receptor tyrosine kinase class V proteins.	BL00790R 16.20 7.677e-09 29-73
207	BL00211	ABC transporters family proteins.	BL00211B 13.37 3.077e-17 573-167
	i	Tarrest Parametria	BL00211B 13.37 7.577e-17 1204-1674

Table 3

SEQ ID NO:	Database entry ID	Description	*Results
			BL00211A 12.23 1.900e-09 472-484
207	PR00478	PHOSPHORIBULOKINASE FAMILY SIGNATURE	PR00478A 13.44 4.133e-09 474-492
207	PR00802	SERUM ALBUMIN FAMILY SIGNATURE	PR00802G 14.57 7.188e-09 971-994
207	PR00836	SOMATOTROPIN HORMONE FAMILY SIGNATURE	PR00836D 13.05 7.125e-09 1504-1519
209	PR00049	WILM'S TUMOUR PROTEIN SIGNATURE	PR00049D 0.00 1.786e-10 288-303
210	BL00972	Ubiquitin carboxyl-terminal hydrolases family 2 proteins.	BL00972D 22.55 3.348e-11 388-413 BL00972E 20.72 4.343e-09 415-437
210	PR00198	ANNEXIN TYPE II SIGNATURE	PR00198H 12.05 7.750e-09 682-696
214	PD00469	PROTEIN PRECURSOR SIGNAL HYDROLA.	PD00469A 13.95 6.400e-09 73-86
215	PF00023	Ank repeat proteins.	PF00023A 16.03 8.875e-10 839-855 PF00023A 16.03 2.286e-09 884-900
215	PR00342	RHESUS BLOOD GROUP PROTEIN SIGNATURE	PR00342H 7.61 9.703e-09 317-340
217	BL00982	Bacterial-type phytoene dehydrogenase proteins.	BL00982A 18.41 8.013e-12 328-360
217	PR00368	FAD-DEPENDENT PYRIDINE NUCLEOTIDE REDUCTASE SIGNATURE	PR00368C 15.74 8.962e-11 326-352
217	PR00469	PYRIDINE NUCLEOTIDE DISULPHIDE REDUCTASE CLASS- II SIGNATURE	PR00469I 13.83 7.532e-11 449-468 PR00469F 16.51 7.152e-09 322-347
217	PD02042	IRON-SULFUR ELECTRON TRANSPORT AROMATIC HYDROCARB.	PD02042B 16.75 5.673e-09 126-141 PD02042A 21.13 9.045e-09 93-120
217	PR00419	ADRENODOXIN REDUCTASE FAMILY SIGNATURE	PR00419A 14.89 9.486e-09 326-349 PR00419D 10.62 9.534e-09 327-342
218	PF00157	PDZ domain proteins (Also known as DHR or GLGF).	PF00157 13.40 4.600e-09 688-699
219	BL00107	Protein kinases ATP-binding region proteins.	BL00107A 18.39 7.000e-23 65-96 BL00107B 13.31 4.214e-10 130-146
219	PR00109	TYROSINE KINASE CATALYTIC DOMAIN SIGNATURE	PR00109B 12.27 7.102e-10 65-84
219	BL00240	Receptor tyrosine kinase class III proteins.	BL00240E 11.56 5.029e-09 51-89
220	PR00239	MOLLUSCAN RHODOPSIN C- TERMINAL TAIL SIGNATURE	PR00239E 1.58 3.045e-09 38-50
220	DM01803	1 HERPESVIRUS GLYCOPROTEIN H.	DM01803A 10.51 9.349e-09 34-55
220	PR00049	WILM'S TUMOUR PROTEIN SIGNATURE	PR00049D 0.00 5.160e-11 40-55 PR00049D 0.00 7.807e-11 41-56 PR00049D 0.00 8.336e-11 38-53 PR00049D 0.00 2.286e-10 42-57 PR00049D 0.00 8.857e-10 33-48 PR00049D 0.00 2.983e-09 37-52 PR00049D 0.00 9.847e-09 43-58
222	BL00326	Tropomyosins proteins.	BL00326A 14.01 5.337e-10 825-859
			220022011 17:01 2:35 /0"10 023-037

Table 3

SEQ ID	Database	Description	*Results
NO:	entry ID		
222	BL01160	Kinesin light chain repeat proteins.	BL01160B 19.54 9.924e-09 516-132
224	BL00478	LIM domain proteins.	BL00478B 14.79 8.527e-09 143-158
226	BL00048	Protamine P1 proteins.	BL00048 6.39 6.063e-09 199-226
228	BL00115	Eukaryotic RNA polymerase II	BL00115Z 3.12 5.744e-10 113-162
		heptapeptide repeat proteins.	BL00115Z 3.12 3.449e-09 120-169
228	BL00415	Synapsins proteins.	BL00415Q 2.23 8.723e-09 253-289
229	BL01161	Glucosamine/galactosamine-6-	BL01161A 19.47 1.000e-40 37-77
	1	phosphate isomerases proteins.	BL01161D 28.14 1.000e-40 199-244
	()	· ·	BL01161B 21.37 5.091e-39 117-160
			BL01161C 18.47 1.500e-23 170-199
231	PR00269	PLEIOTROPHIN/MIDKINE FAMILY SIGNATURE	PR00269A 13.91 3.133e-30 88-113
231	BL00181	PTN/MK heparin-binding protein	BL00181A 19.07 4.960e-37 76-112
		family proteins.	BL00181A 19:07 9.224e-18 78-114
236	BL00888	Cyclic nucleotide-binding domain proteins.	BL00888B 14.79 9.069e-13 499-523
236	BL00415	Synapsins proteins.	BL00415N 4.29 2.774e-09 733-777
236	PD00306	PROTEIN GLYCOPROTEIN PRECURSOR RE.	PD00306A 10.26 3.133e-09 646-660
236	PR00209	ALPHA/BETA GLIADIN FAMILY SIGNATURE	PR00209B 4.88 3.813e-09 739-758
236	DM00668	ZEIN.	DM00668A 10.20 8.500e-09 258-273
238	BL01188	GNS1/SUR4 family proteins.	BL01188B 13.46 4.115e-26 120-151
			BL01188C 22.65 4.136e-26 151-202
	ļ		BL01188D 8.62 1.290e-11 238-255
			BL01188A 18.82 6.718e-10 55-87
239	PR00929	AT-HOOK-LIKE DOMAIN	PR00929B 4.38 8.875e-09 133-583
		SIGNATURE	PR00929C 5.26 8.914e-09 133-144
242	BL00232	Cadherins extracellular repeat proteins	BL00232B 32.79 2.765e-25 541-151
		domain proteins.	BL00232B 32.79 8.263e-22 766-814
			BL00232B 32.79 2.397e-21 67-115
			BL00232B 32.79 4.133e-19 1481-1529
		•	BL00232B 32.79 1.000e-18 1371-1419
			BL00232B 32.79 2.662e- 18 1691-1739
			BL00232B 32.79 5.292e-18 1287-1335
			BL00232B 32.79 9.147e-18 1148-1196
			BL00232B 32.79 1.265e-17 980-1028
			BL00232B 32.79 1.529e-17 426-474
		.*	BL00232B 32.79 2.588e-17 1084-1132
·		·	BL00232B 32.79 1.386e-16 1184-1232
			BL00232C 10.65 5.390e-12 1369-1387
		·	BL00232C 10.65 1.391e-11 204-660
			BL00232C 10.65 2.174e-11 1584-1164 BL00232C 10.65 4.522e-11 1689-1707
		· .	BL00232C 10.65 1.000e-10 65-83 BL00232C 10.65 4.115e-10 1285-1303
			BL00232C 10.65 4.115e-10 1285-1303 BL00232B 32.79 7.200e-10 649-697
			BL00232B 32.79 7.200e-10 649-697 BL00232C 10.65 9.827e-10 978-996
			BL00232C 10.65 9.827e-10 978-996 BL00232C 10.65 1.947e-09 170-188
	· .		BL00232C 10.65 1.947e-09 170-188 BL00232B 32.79 2.137e-09 172-220
	li		DLNU434D 34.17 4.13 /6-07 1 /4-440

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Table 3

SEQ ID NO:	Database entry ID	Description	*Results
110.	CILC. Y ALD		BL00232C 10.65 4.474e-09 1182-1200
			BL00232C 10.65 8.737e-09 539-119
243	BL00795	Involucrin proteins.	BL00795C 17.06 4.977e-10 64-109
			BL00795C 17.06 6.300e-09 55-100
244	BL00790	Receptor tyrosine kinase class V	BL00790I 20.01 7.823e-15 23-54 BL00790I
•	·	proteins.	20.01 9.400e-11 310-341 BL00790I 20.01
	1		1.900e-10 117-148 BL00790I 20.01 3.893e-
		· · · · · · · · · · · · · · · · · · ·	09 215-246
244	PR00014	FIBRONECTIN TYPE III REPEAT	PR00014D 12.04 6.400e-11 30-45
	•	SIGNATURE	PR00014D 12.04 6.400e-11 317-332
	1	·	PR00014C 15.44 9.171e-09 204-223
2.12			PR00014D 12.04 1.000e-08 222-237
245	BL00183	Ubiquitin-conjugating enzymes proteins.	BL00183 28.97 7.037e-10 140-188
246	PR00019	LEUCINE-RICH REPEAT	PR00019A 11.19 8.800e-12 205-219
		SIGNATURE	PR00019B 11.36 2.000e-11 202-216
247	BL00214	Cytosolic fatty-acid binding proteins.	BL00214B 26.51 7.180e-24 206-251
		• .	BL00214A 21.17 6.250e-22 165-191
247	PR00178	FATTY ACID-BINDING PROTEIN	PR00178A 15.07 4.913e-21 166-187
	!	SIGNATURE	PR00178C 20.54 2.500e-17 226-254
			PR00178D 13.52 6.897e-16 272-291
2.12	 		PR00178B 10.52 4.900e-10 200-212
248	PR00395	RIBOSOMAL PROTEIN S2 SIGNATURE	PR00395C 16.17 2.047e-13 46-64
248	BL00962	Ribosomal protein S2 proteins.	BL00962C 15.90 2.846e-12 46-64
249	BL00227	Tubulin subunits alpha, beta, and	BL00227D 18.46 1.000e-40 74-128
		gamma proteins.	BL00227F 21.16 1.529e-33 226-280
			BL00227E 24.15 1.409e-26 178-213
250	BL00227	Tubulin subunits alpha, beta, and	BL00227C 25.48 1.000e-40 39-91
	. '	gamma proteins.	BL00227D 18.46 1.000e-40 148-202
	1		BL00227F 21.16 1.529e-33 300-354
200			BL00227E 24.15 1.409e-26 252-287
251	BL00152	ATP synthase alpha and beta subunits	BL00152B 21.40 1.900e-31 191-229
	[proteins.	BL00152A 15.38 5.154e-21 134-160
250	DT 00150	ATTO	BL00152C 11.41 6.250e-12 291-303
252	BL00152	ATP synthase alpha and beta subunits	BL00152E 22.68 1.000e-32 285-323
	!	proteins.	BL00152A 15.38 5.154e-21 134-160 BL00152C 11.41 6.250e-12 247-259
253	BL00518	Zinc finger, C3HC4 type (RING	BL00518 12.23 2.200e-11 54-63
255	PLU0219	finger), proteins.	BL00316 12.23 2.2006-11 34-03
253	BL01282	BIR repeat proteins.	BL01282B 30,49 2.029e-09 35-74
254	DM00892	3 RETROVIRAL PROTEINASE.	DM00892C 23.55 9.739e-12 417-451
254	PR00417	PROKARYOTIC DNA	PR00417A 12.66 8.472e-09 65-79
		TOPOISOMERASE I SIGNATURE	
255	BL01052	Calponin family repeat proteins.	BL01052C 18.51 1.000e-40 88-128
	1		BL01052A 16.12 2.875e-35 3-35 BL01052B
	 		15.31 5.219e-26 52-78
255	PR00888	SMOOTH MUSCLE	PR00888D 16.09 9.112e-19 89-106
	}	PROTEIN/CALPONIN FAMILY	PR00888E 11.81 2.800e-18 105-121
	<u> </u>	SIGNATURE	PR00888F 7.44 4.600e-18 126-141

Table 3

SEQ ID NO:	Database entry ID	Description	*Results
			PR00888A 11.87 7.750e-18 3-18 PR00888C 12.27 2.286e-17 52-68 PR00888G 12.73 9.438e-15 163-177 PR00888B 13.72 1.321e-
255	DDOOGO	CN 40 COTT N 41 10 CUE PROTERIA CO	14 22-36
255	PR00890	SMOOTH MUSCLE PROTEIN 22- ALPHA (TRANSGELIN)	PR00890E 14.34 1.429e-27 136-156 PR00890A 8.61 1.000e-26 34-54 PR00890C
-		SIGNATURE	8.22 1.600e-19 85-99 PR00890B 8.75
		BIGIVITORE	6.318e-19 62-78 PR00890F 12.92 1.205e-17
			162-175 PR00890D 16.17 1.130e-13 119-
	l i		129
257	BL00745	Prokaryotic-type class I peptide chain	BL00745C 13.66 1.000e-40 202-249
		release factors signat.	BL00745B 22.56 8.683e-33 148-191
			BL00745D 14.90 8.435e-23 280-303
259	BL00194	Thioredoxin family proteins.	BL00194 12.16 7.429e-10 684-697
260	BL00612	Osteonectin domain proteins.	BL00612E 13.12 3.948e-10 391-436
260	BL00484	Thyroglobulin type-1 repeat proteins	BL00484C 17.01 8.244e-11 136-151
	[proteins.	BL00484B 9.04 2.145e-10 249-263
	1		BL00484C 17.01 2.309e-09 269-284 BL00484B 9.04 8.950e-09 116-130
262	PR00187	DNAJ PROTEIN FAMILY	PR00187A 12.84 2.375e-09 288-308
202	1100187	SIGNATURE	FR00187A 12.84 2.5736-03 288-308
262	BL00198	Nt-dnaJ domain proteins.	BL00198A 8.07 3.681e-09 292-309
262	BL00157	Aminotransferases class-V pyridoxal-	BL00157A 11.72 8.200e-09 16-26
		phosphate attachment site proteins.	
263	PR00320	G-PROTEIN BETA WD-40 REPEAT SIGNATURE	PR00320B 12.19 2.125e-09 207-222
263	PF00913	Trypanosome variant surface glycoprotein.	PF00913A 7.33 2.500e-09 666-673
266	BL01144	Ribosomal protein L31e proteins.	BL01144 25.07 1.000e-40 21-73
268	DM00516	186 DISCOIDIN I N-TERMINAL.	DM00516 30.53 8.168e-13 153-198
268	BL00132	Zinc carboxypeptidases, zinc-binding	BL00132C 21.35 7.863e-10 307-348
<u> </u>		region 1 proteins.	BL00132A 26.07 8.988e-10 224-265
268	PR00765	CARBOXYPEPTIDASE A	PR00765B 15.57 7.171e-12 276-291
	}	METALLOPROTEASE (M14)	PR00765D 14.16 1.551e-09 420-434
268	BL00170	FAMILY SIGNATURE Cyclophilin-type peptidyl-prolyl cis-	BL00170A 17.08 9.018e-09 485-512
200	BLOOT	trans isomerase signatur.	BL00170A 17.08 9.018e-09 485-512
269	BL00622	Bacterial regulatory proteins, luxR	BL00622 32.69 9.780e-09 11-58
		family proteins.	·
270	PR00048	C2H2-TYPE ZINC FINGER	PR00048A 10.52 1.000e-11 447-461
	}	SIGNATURE	PR00048A 10.52 4.316e-11 389-403
			PR00048A 10.52 6.684e-11 362-376
270	PF00651	BTB (also known as BR-C/Ttk) domain proteins.	PF00651 15.00 3.143e-10 37-50
270	BL00028	Zinc finger, C2H2 type, domain	BL00028 16.07 7.000e-10 392-409 BL00028
	j	proteins.	16.07 9.100e-10 256-273 BL00028 16.07
			2.286e-09 450-467 BL00028 16.07 8.714e-
074	722 (00000	CYPA 11 YOUR DESCRIPTION	09 365-382
274	DM00303	6 LEA 11-MER REPEAT REPEAT.	DM00303A 13.20 3.310e-09 467-517
275	PF00622	Domain in SPla and the RYanodine	PF00622B 21.00 9.357e-14 374-396

Table 3

SEQ ID NO:	Database entry ID	Description	*Results
		Receptor.	PF00622C 12.62 1.857e-12 458-472
275	BL00518	Zinc finger, C3HC4 type (RING	BL00518 12.23 8.800e-11 44-53
	<u> </u>	finger), proteins.	
277	PF00651	BTB (also known as BR-C/Ttk) domain	PF00651 15.00 9.133e-10 65-78
		proteins.	
278	PD00066	PROTEIN ZINC-FINGER METAL-	PD00066 13.92 8.200e-16 295-308 PD00066
	l	BINDI.	13.92 8.200e-16 519-532 PD00066 13.92
	Í		1.692e-15 351-364 PD00066 13.92 4.462e-
			15 547-122 PD00066 13.92 4.600e-14 323-
	l	,	336 PD00066 13.92 4.600e-14 435-448
	ì		PD00066 13.92 7.000e-14 463-476 PD00066
			13.92 1.500e-13 239-252 PD00066 13.92 3.143e-12 267-280 PD00066 13.92 3.143e-
		<u> </u>	12 407-420 PD00066 13.92 8.826e-11 211-
		·	224 PD00066 13.92 2.038e-10 491-504
		·	PD00066 13.92 2.385e-10 379-392
278	PR00048	C2H2-TYPE ZINC FINGER	PR00048A 10.52 7.750e-16 444-458
	1	SIGNATURE	PR00048A 10.52 6.727e-15 360-374
	}		PR00048A 10.52 9.182e-15 528-542
			PR00048A 10.52 7.000e-14 472-486
			PR00048A 10.52 7.750e-14 388-402
		•	PR00048A 10.52 1.000e-13 332-346
			PR00048A 10.52 3.133e-13 304-318
			PR00048A 10.52 4.857e-13 118-132
		·	PR00048A 10.52 6.786e-13 500-514
			PR00048B 6.02 1.000e-12 292-302
			PR00048A 10.52 8.941e-12 192-206
			PR00048B 6.02 1.000e-11 348-358
		·	PR00048A 10.52 1.947e-11 248-262
		,	PR00048B 6.02 2.385e-11 264-274
,	·		PR00048B 6.02 7.231e-11 544-116
			PR00048A 10.52 7.632e-11 416-430
			PR00048B 6.02 8.615e-11 236-246
			PR00048B 6.02 2.688e-10 516-526
			PR00048B 6.02 4.375e-10 460-470 PR00048B 6.02 4.375e-10 488-498
			PR00048B 6.02 4.5756-10 466-496
			PR00048B 6.02 6.063e-10 320-330
			PR00048A 10.52 7.214e-10 220-234
			PR00048B 6.02 1.947e-09 432-442
		·	PR00048B 6.02 4.316e-09 572-144
278	DM01970	0 kw ZK632.12 YDR313C	DM01970B 8.60 5.012e-09 191-204
		ENDOSOMAL III.	
279	PD00066	PROTEIN ZINC-FINGER METAL-	PD00066 13.92 6.400e-16 449-462 PD00066
		BINDI.	13.92 6.538e-15 504-517 PD00066 13.92
			9.308e-15 421-434 PD00066 13.92 7.000e-
			14 476-489 PD00066 13.92 6.087e-11 393-
			406
279	BL00028	Zinc finger, C2H2 type, domain	BL00028 16.07 2.500e-17 350-367 BL00028
		proteins.	16.07 5.050e-13 405-422 BL00028 16.07

Table 3

SEQ ID	Database	Description	*Results
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			9.171e-12 433-450 BL00028 16.07 2.731e-
	i		488-505 BL00028 16.07 3.077e-11 516-533
	1		BL00028 16.07 6.100e-10 377-394
279	PD02462	PROTEIN BOLA TRANSCRIPTION REGULATION AC.	PD02462A 22.48 6.488e-09 481-516
279	PR00048	C2H2-TYPE ZINC FINGER	PR00048A 10.52 3.250e-16 347-361
		SIGNATURE	PR00048B 6.02 5.154e-11 501-511
	1.		PR00048B 6.02 1.000e-10 446-456
			PR00048A 10.52 1.391e-10 513-527
			PR00048A 10.52 2.565e-10 485-499
			PR00048A 10.52 5.696e-10 402-416
		v .	PR00048B 6.02 8.875e-10 418-428
			PR00048A 10.52 1.720e-09 430-444
	1	•	PR00048B 6.02 3.368e-09 390-400
	ļ		PR00048A 10.52 8.200e-09 374-388
285	BL00276	Channel forming colicins proteins.	BL00276A 8.87 6.500e-09 257-269
286	PD01066	PROTEIN ZINC FINGER ZINC-	PD01066 19.43 2.000e-30 10-49
		FINGER METAL-BINDING NU.	1201000 13.43 2.0000-30 10-43
286	PD00066	PROTEIN ZINC-FINGER METAL-	PD00066 13.92 6.400e-16 388-401 PD00066
		BINDI.	13.92 3.769e-15 248-261 PD00066 13.92
			9.308e-15 304-317 PD00066 13.92 2.200e-
			14 360-373 PD00066 13.92 2,200e-14 416-
			429 PD00066 13.92 6.400e-14 332-345
	1		PD00066 13.92 1.000e-13 220-233 PD00066
• •	<u> </u>		13.92 2.500e-13 192-205 PD00066 13.92
			5.000e-13 276-289 PD00066 13.92 5.500e-
		·	09 136-149
286	BL00028	Zinc finger, C2H2 type, domain	BL00028 16.07 2.286e-16 260-277 BL00028
	ļ ·	proteins.	16.07 2.588e-14 288-305 BL00028 16.07
	1	[• ·	2.800e-13 400-417 BL00028 16.07 6.850e-
,			13 120-137 BL00028 16.07 3.423e-11 148-
	ŀ	·	165 BL00028 16.07 7.923e-11 344-361
			BL00028 16.07 2.500e-10 204-221 BL00028
			16.07 2.500e-10 428-445 BL00028 16.07
	1		3.100e-10 316-333 BL00028 16.07 6.100e-
			10 176-193 BL00028 16.07 1.771e-09 232-
			249 BL00028 16.07 8.200e-09 372-389
286	PR00048	C2H2-TYPE ZINC FINGER	PR00048A 10.52 7.000e-17 257-271
	}	SIGNATURE	PR00048A 10.52 6.727e-15 397-411
		• •	PR00048A 10.52 2.929e-13 285-299
		· · · · · · · · · · · · · · · · · · ·	PR00048A 10.52 9.471e-12 369-383
	1 1		PR00048B 6.02 1.000e-11 329-339
]		PR00048A 10.52 1.474e-11 313-327
			PR00048A 10.52 2.421e-11 425-439
	1		PR00048B 6.02 3.077e-11 385-395
	1		PR00048A 10.52 6.684e-11 117-131
			PR00048A 10.52 8.141e-11 201-215
	[.		PR00048A 10.52 1.783e-10 341-355
	<u> </u>	<u> </u>	PR00048B 6.02 2.125e-10 301-311

Table 3

SEQ ID NO:	Database entry ID	Description	*Results
			PR00048B 6.02 2.125e-10 357-367
	}		PR00048B 6.02 2.688e-10 217-227
			PR00048A 10.52 3.739e-10 229-243
			PR00048B 6.02 4.938e-10 273-283
	l .		PR00048B 6.02 1.474e-09 245-255
•			PR00048A 10.52 2.440e-09 145-159
	1		PR00048B 6.02 3.842e-09 161-171
		•	PR00048B 6.02 8.105e-09 441-451
	<u> </u>		PR00048B 6.02 9.053e-09 189-199
287	PD01066	PROTEIN ZINC FINGER ZINC- FINGER METAL-BINDING NU.	PD01066 19.43 7.407e-23 3-42
287	BL00028	Zinc finger, C2H2 type, domain	BL00028 16.07 8.941e-14 269-286 BL00028
201	1000020	proteins.	16.07 1.000e-13 549-128 BL00028 16.07
•	l	process.	2.565e-12 194-650 BL00028 16.07 6.087e-
]	. 11	12 241-258 BL00028 16.07 6.870e-12 297-
,		,	314 BL00028 16.07 6.870e-12 381-398
1	[BL00028 16.07 7.214e-12 493-510 BL00028
	}		16.07 1.346e-11 465-482 BL00028 16.07
			1.692e-11 353-370 BL00028 16.07 3.769e-
			11 325-342 BL00028 16.07 6.192e-11 167-
			622 BL00028 16.07 8.962e-11 213-230
			BL00028 16.07 1.600e-10 409-426 BL00028
			16.07 5.200e-10 185-202 BL00028 16.07
	[6.700e-10 577-156 BL00028 16.07 3.057e-
			09 521-538 BL00028 16.07 6.143e-09 437-
	j		454
287	PR00048	C2H2-TYPE ZINC FINGER	PR00048A 10.52 9.250e-14 238-252
		SIGNATURE	PR00048A 10.52 3.209e-12 266-280
			PR00048A 10.52 4.706e-12 490-504
•			PR00048A 10.52 5.765e-12 462-476
			PR00048A 10.52 7.882e-12 630-644
			PR00048A 10.52 8.941e-12 518-532
			PR00048A 10.52 9.471e-12 164-178
]		PR00048A 10.52 5.737e-11 378-392
	1		PR00048A 10.52 7.158e-11 546-122
,	.		PR00048B 6.02 7.231e-11 180-190
			PR00048A 10.52 8.141e-11 210-224
			PR00048A 10.52 9.053e-11 294-308
		•	PR00048A 10.52 9.053e-11 406-420
			PR00048A 10.52 3.348e-10 322-336
		,	PR00048B 6.02 3.813e-10 338-348
	(·		PR00048B 6.02 3.813e-10 394-404
	(PR00048B 6.02 3.813e-10 478-488
]		PR00048B 6.02 4.938e-10 506-516
,]		PR00048A 10.52 8.043e-10 434-448
١.			PR00048B 6.02 8.875e-10 226-236
]		PR00048B 6.02 8.875e-10 450-460
]		PR00048B 6.02 1.000e-09 366-376
!			PR00048B 6.02 1.000e-09 422-432
			PR00048A 10.52 3.520e-09 136-588

Table 3

SEQ ID NO:	Database entry ID	Description	*Results
			PR00048B 6.02 7.158e-09 590-600
			PR00048B 6.02 7.632e-09 310-320
			PR00048B 6.02 7.632e-09 124-572
			PR00048A 10.52 9.280e-09 350-364
289	PR00070	DIHYDROFOLATE REDUCTASE	PR00070C 13.09 6.143e-16 51-63
		SIGNATURE	PR00070D 11.63 2.929e-15 112-127
289	BL00075	Dihydrofolate reductase proteins.	BL00075A 27.70 7.900e-16 8-39 BL00075B
	ļ		13.49 3.813e-15 51-63 BL00075C 8.51
:			2.862e-11 66-79 BL00075D 5.74 8.105e-10
		<u></u>	113-123
292	PR00250	FUNGAL PHEROMONE MATING	PR00250D 14.62 9.163e-09 254-278
	L	FACTOR STE2 GPCR SIGNATURE	
294	PR00081	GLUCOSE/RIBITOL	PR00081A 10.53 2.731e-09 39-57
	'	DEHYDROGENASE FAMILY	
	<u> </u>	SIGNATURE	
294	PR00080	ALCOHOL DEHYDROGENASE	PR00080C 17.16 6.464e-11 191-211
		SUPERFAMILY SIGNATURE	PR00080A 9.32 9.750e-09 118-130
295	PR00806	VINCULIN SIGNATURE	PR00806B 4.28 8.920e-09 276-290
		,	PR00806B 4.28 9.202e-09 275-289
296	PF00992	Troponin.	PF00992A 16.67 3.789e-10 553-588
296	BL00752	XPA protein.	BL00752B 19.17 8.144e-09 130-612
296	BL01160	Kinesin light chain repeat proteins.	BL01160B 19.54 8.551e-09 536-590
298	PR00511	TEKTIN SIGNATURE	PR00511C 7.86 4.214e-09 371-388
300	BL00353	HMG1/2 proteins.	BL00353B 11.47 9.171e-19 228-278
301	PR00240	ALPHA-1A ADRENERGIC RECEPTOR SIGNATURE	PR00240C 8.38 3.941e-10 316-336
302	BL00518	Zinc finger, C3HC4 type (RING finger), proteins.	BL00518 12.23 2.200e-11 54-63
302	BL01282	BIR repeat proteins.	BL01282B 30.49 2.029e-09 35-74
305	PR00193	MYOSIN HEAVY CHAIN	PR00193D 14.36 1.545e-31 390-419
		SIGNATURE	PR00193C 12.60 1.209e-25 143-171
	Į.		PR00193B 11.69 2.543e-24 95-121
	1	·	PR00193A 15.41 6.885e-19 39-59
			PR00193E 19.47 3.291e-12 444-473
305	BL00675	Sigma-54 interaction domain proteins	BL00675A 24.86 3.475e-09 98-142
	1	ATP-binding region A proteins.	
306	PR00239	MOLLUSCAN RHODOPSIN C-	PR00239E 1.58 5.920e-11 47-59
	i ·	TERMINAL TAIL SIGNATURE	
306	PD00066	PROTEIN ZINC-FINGER METAL-	PD00066 13.92 7.923e-15 140-153 PD00066
•	[BINDI.	13.92 4.000e-14 112-125 PD00066 13.92
			1.391e-11 84-97 PD00066 13.92 1.692e-10
			168-181
306	BL00028	Zinc finger, C2H2 type, domain	BL00028 16.07 2.059e-14 96-113 BL00028
		proteins.	16.07 4.130e-12 124-141 BL00028 16.07
	1	·.	2.385e-11 68-85 BL00028 16.07 8.269e-11
			180-197 BL00028 16.07 8.962e-11 152-169
			BL00028 16.07 9.400e-10 319-336
306	PR00799	ASPARTATE	PR00799D 16.46 5.125e-09 188-214
		AMINOTRANSFERASE SIGNATURE	A TOTAL TOTAL OF THE STATE OF T

Table 3

SEQ ID	Database	Description	*Results
NO:	entry ID	Protect	ALGUID
306	PR00048	C2H2-TYPE ZINC FINGER	PR00048B 6.02 1.900e-13 81-91 PR00048A
	1	SIGNATURE	10.52 3.133e-13 65-79 PR00048A 10.52
	İ	i i	9.357e-13 121-135 PR00048A 10.52 9.357e-
			13 149-163 PR00048B 6.02 2.688e-10 137-
	1		147 PR00048A 10.52 4.522e-10 279-293
			PR00048A 10.52 5.696e-10 177-191
	ì		PR00048B 6.02 9.438e-10 109-119
	1		PR00048A 10.52 3.160e-09 93-107
1			PR00048B 6.02 8.105e-09 165-175
307	PD00015	GLYCOPROTEIN PRECURSOR	PD00015A 8.90 6.400e-09 35-43
	ł	CELL SI.	
310	DM00031	IMMUNOGLOBULIN V REGION.	DM00031B 15.41 3.662e-11 80-114
311	BL00824	Elongation factor 1 beta/beta/delta	BL00824C 14.58 1.000e-40 129-167
	ļ	chain proteins.	BL00824D 14.04 6.192e-39 167-202
	<u> </u>	· .	BL00824B 9.21 2.080e-21 96-116
	<u>.</u>		BL00824E 12.49 3.333e-19 210-226
312	PR00501	KELCH REPEAT SIGNATURE	PR00501B 18.88 7.632e-09 476-491
	ļ		PR00501B 18.88 9.763e-09 523-538
313	PD01066	PROTEIN ZINC FINGER ZINC-	PD01066 19.43 6.200e-30 43-82
		FINGER METAL-BINDING NU.	
313	PD00066	PROTEIN ZINC-FINGER METAL	PD00066 13.92 6.500e-13 439-452 PD00066
		BINDI	13.92 8.000e-13 355-368 PD00066 13.92
			1.000e-12 383-396 PD00066 13.92 4.000e-
			12 327-340 PD00066 13.92 5.714e-12 411-
		•	424 PD00066 13.92 8.435e-11 299-31213.92
			5.800e-14 467-480 PD00066
313	BL00028	Zinc finger, C2H2 type, domain	BL00028 16.07 2.565e-12 451-468 BL00028
		proteins.	16.07 2.957e-12 311-328 BL00028 16.07
			3.348e-12 367-384 BL00028 16.07 1.692e-
		•	11 423-440 BL00028 16.07 2.731e-11 283-
		•	300 BL00028 16.07 2.800e-10 339-356
		,	BL00028 16.07 9.700e-10 199-216 BL00028
	·	•	16.07 1.000e-09 395-412 BL00028 16.07
			4.086e-09 120-137
313	PR00048	C2H2-TYPE ZINC FINGER	PR00048A 10.52 5.909e-15 364-378
] . ;	SIGNATURE	PR00048A 10.52 2.286e-13 308-322
İ	1		PR00048A 10.52 7.429e-13 392-406
٠.	!		PR00048A 10.52 6.824e-12 448-462
			PR00048A 10.52 2.421e-11 196-210
			PR00048A 10.52 1.000e-10 280-294
			PR00048B 6.02 3.813e-10 324-334
	· ·		PR00048B 6.02 4.375e-10 464-474
			PR00048A 10.52 6.870e-10 336-350
	[PR00048A 10.52 7.214e-10 420-434
			PR00048B 6.02 7.750e-10 436-446
-			PR00048B 6.02 4.316e-09 380-390
314	PR00121	SODIUM/POTASSIUM-	PR00121D 16.72 1.577e-13 210-232
		TRANSPORTING ATPASE	
		SIGNATURE	<u> </u>
314	PR00119	P-TYPE CATION-TRANSPORTING	PR00119B 13.94 9.194e-12 217-232

Table 3

SEQ ID	Database	Description	*Results
NO:	entry ID		
		ATPASE SUPERFAMILY SIGNATURE	
314	BL01228	Hypothetical cof family proteins.	BL01228D 17.44 3.400e-11 646-671
314	BL00154	E1-E2 ATPases phosphorylation site	BL00154E 20.37 4.054e-13 486-527
		proteins.	BL00154C 12.38 4.060e-12 213-232
		:	BL00154F 8.23 9.597e-11 207-669
315	BL00888	Cyclic nucleotide-binding domain proteins.	BL00888B 14.79 1.692e-10 396-420
315	BL00420	Speract receptor repeat proteins domain proteins.	BL00420A 20.42 8.338e-09 215-682
315	DM00668	ZEIN.	DM00668A 10.20 8.500e-09 155-170
316	PR00727	BACTERIAL LEADER PEPTIDASE 1	PR00727C 13.04 9.063e-16 108-128
		(S26) FAMILY SIGNATURE	PR00727B 12.51 7.848e-11 81-94
316	BL00501	Signal peptidases I serine proteins.	BL00501D 16.69 2.884e-13 108-128
		- -	BL00501C 9.61 9.561e-11 81-93 BL00501B
			12.58 7.000e-09 61-77
317	PD01066	PROTEIN ZINC FINGER ZINC- FINGER METAL-BINDING NU.	PD01066 19.43 9.471e-27 13-52
317	BL00028	Zinc finger, C2H2 type, domain	BL00028 16.07 5.235e-14 214-231 BL00028
		proteins.	16.07 6.850e-13 270-287 BL00028 16.07
	ĺ	÷	9.100e-13 354-371 BL00028 16.07 1.391e-
	1		12 158-175 BL00028 16.07 1.346e-11 298-
			315 BL00028 16.07 3.769e-11 242-259
		· ·	BL00028 16.07 6.538e-11 380-397 BL00028
			16.07 8.800e-10 186-203 BL00028 16.07
			1.514e-09 326-343
317	PR00048	C2H2-TYPE ZINC FINGER	PR00048B 6.02 3.000e-12 199-209
		SIGNATURE	PR00048A 10.52 7.882e-12 351-365
			PR00048A 10.52 8.412e-12 323-337
		•	PR00048A 10.52 8.941e-12 239-253
			PR00048A 10.52 1.474e-11 211-225
			PR00048A 10.52 6.211e-11 155-169
		·	PR00048B 6.02 7.231e-11 311-321
			PR00048A 10.52 8.141e-11 267-281
			PR00048B 6.02 3.250e-10 339-349
			PR00048B 6.02 3.813e-10 255-265
			PR00048B 6.02 7.188e-10 283-293
	,		PR00048B 6.02 3.842e-09 171-181
			PR00048B 6.02 3.842e-09 393-403
319	PR00004	ANA DYDA ATOVDI DOLLARI	PR00048A 10.52 8.200e-09 295-309
		ANAPHYLATOXIN DOMAIN SIGNATURE	PR00004C 12.46 8.141e-09 91-103
320	DM00060	338 kw NEUREXIN ALPHA III CYSTEINE.	DM00060 6.92 6.500e-11 28-38
320	PR00010	TYPE II EGF-LIKE SIGNATURE	PR00010C 11.16 7.667e-11 44-55
325	PR00020	MAM DOMAIN SIGNATURE	PR00020A 18.17 5.776e-12 344-363
			PR00020C 13.66 6.932e-10 417-429
325	BL00740	MAM domain proteins.	BL00740A 13.87 8.313e-12 346-359
		· ·	BL00740B 19.76 8.500e-09 486-507
325	PD02080	T-CELL GLYCOPROTEIN CD8	PD02080B 20.69 9.621e-09 123-162

Table 3

SEQ ID NO:	Database entry ID	Description	*Results
		CHAIN SURFACE ALPHA PRE.	
326	BL00048	Protamine P1 proteins.	BL00048 6.39 6.128e-10 167-194
326	PF01140	Matrix protein (MA), p15.	PF01140D 15.54 9.791e-09 220-255
327	PR00020	MAM DOMAIN SIGNATURE	PR00020C 13.66 2.615e-11 143-593
			PR00020B 15.52 5.059e-10 52-69
	1		PR00020B 15.52 1.789e-09 553-132
329	PD01066	PROTEIN ZINC FINGER ZINC-	PD01066 19.43 9.357e-32 8-47
	1	FINGER METAL-BINDING NU.	
329	BL00028	Zinc finger, C2H2 type, domain	BL00028 16.07 3.209e-14 284-301 BL00028
		proteins.	16.07 4.600e-13 508-525 BL00028 16.07
	i	•	6.400e-13 368-385 BL00028 16.07 4.115e-
	}		11 396-413 BL00028 16.07 4.115e-11 424-
			441 BL00028 16.07 8.269e-11 172-189
			BL00028 16.07 8.962e-11 256-273 BL00028
		<u>.</u>	16.07 9.308e-11 312-329 BL00028 16.07
			9.654e-11 200-217 BL00028 16.07 3.100e-
		,	10 340-357 BL00028 16.07 5.500e-10 452-
			469 BL00028 16.07 9.100e-10 480-497
]	<u> </u>	BL00028 16.07 4.086e-09 228-245
329	PD00066	PROTEIN ZINC-FINGER METAL-	PD00066 13.92 7.000e-14 272-285 PD00066
		BINDI.	13.92 5.000e-13 328-341 PD00066 13.92
			5.500e-13 188-201 PD00066 13.92 5.500e-
	[13 384-397 PD00066 13.92 6,000e-13 496-
			509 PD00066 13.92 6.143e-12 468-481
			PD00066 13.92 2.731e-10 440-453 PD00066
			13.92 4.808e-10 160-173 PD00066 13.92
			5.500e-10 244-257 PD00066 13.92 7.000e-
			09 216-229 PD00066 13.92 7.000e-09 412- 425
332	PD02870	RECEPTOR INTERLEUKIN-1	PD02870B 18.83 5.871e-11 468-501
552	1202070	PRECURSOR.	1 D02870D 18.83 3.8716-11 406-301
332	PR00019	LEUCINE-RICH REPEAT	PR00019A 11.19 8.043e-10 275-289
552	1200015	SIGNATURE	TROUTSA 11.19 8.0436-10 273-289
332	BL00240	Receptor tyrosine kinase class III	BL00240B 24.70 4.447e-09 430-454
	2202.0	proteins.	BE00240B 24.70 4.4470-05 450-454
333	BL00738	S-adenosyl-L-homocysteine hydrolase	BL00738J 18.61 1.000e-40 154-204
		proteins.	BL00738H 23.08 5.320e-36 468-521
		Protonal.	BL00738F 12.23 7.261e-29 387-419
			BL00738A 16.27 9.660e-27 216-256
			BL00738C 16.53 7.923e-25 281-319
			BL00738G 14.29 6.268e-23 446-468
	-		BL00738B 12.28 8.085e-21 256-281
			BL00738E 14.18 9.200e-19 361-384
	,		BL00738I 14.57 5.135e-17 545-583
	'		BL00738D 7.16 5.109e-13 335-350
333	BL00836	Alanine dehydrogenase & pyridine	BL00836D 22.30 8.622e-09 424-461
		nucleotide transhydrogenase.	
337	PR00425	BRADYKININ RECEPTOR	PR00425C 13.23 3.148e-09 80-100
		SIGNATURE	
342	PD01823		PD01823E 9.30 6.824e-12.108-121
342	PD01823	PROTEIN INTERGENIC REGION	PD01823E 9.30 6.824e-12 108-121

Table 3

SEQ ID NO:	Database entry ID	Description	*Results			
		ABC1 PRECURSOR	PD01823D 16.66 1.265e-09 46-67			
	1	MITOCHONDRION T.				
343	PR00976	RIBOSOMAL PROTEIN S21	PR00976C 10.41 2.837e-09 396-407			
242	D) (00015	FAMILY SIGNATURE.	D) (00015 10 40 1 450 00 450 506			
343	DM00215	PROLINE-RICH PROTEIN 3.	DM00215 19.43 1.458e-09 473-506			
242	PROCES	DATE DE CALLEY	DM00215 19.43 4.814e-09 463-496			
343	PR00671	INHIBIN BETA B CHAIN SIGNATURE	PR00671C 4.18 9.172e-09 707-727			
343	PD01234	PROTEIN NUCLEAR	PD01234B 15.53 1.000e-08 482-500			
	<u> </u>	BROMODOMAIN TRANS.				
344	PR00175	MYOGLOBIN SIGNATURE	PR00175B 9.02 2.143e-10 25-49			
344	PR00814	BETA HAEMOGLOBIN SIGNATURE	PR00814C 9.20 6.523e-10 66-84			
344	PR00173	ERYTHROCRUORIN FAMILY SIGNATURE	PR00173A 15.91 7.158e-10 25-48			
344	BL01033	Globins profile.	BL01033A 16.94 1.000e-16 25-47			
			BL01033B 13.81 8.615e-09 87-99			
344	PR00612	ALPHA HAEMOGLOBIN	PR00612E 9.04 4.194e-12 122-139			
		SIGNATURE	PR00612B 10.92 3.483e-10 32-43			
	[PR00612D 9.76 9.438e-09 74-88			
345	PR00814	BETA HAEMOGLOBIN SIGNATURE	PR00814C 9.20 6.523e-10 104-122			
345	BL01033	Globins profile.	BL01033A 16.94 5.125e-10 63-85			
		Cassand promo.	BL01033B 13.81 8.615e-09 125-137			
345	PR00612	ALPHA HAEMOGLOBIN	PR00612E 9.04 4.194e-12 160-177			
		SIGNATURE	PR00612B 10.92 3.483e-10 70-81			
]		PR00612D 9.76 9.438e-09 112-126			
349	PD01066	PROTEIN ZINC FINGER ZINC-	PD01066 19.43 6.133e-32 6-45			
350	DT 00072	FINGER METAL-BINDING NU.	DT 00072 A 11 02 C 210 10 2C4 202			
330	BL00972	Ubiquitin carboxyl-terminal hydrolases	BL00972A 11.93 6.318e-19 364-382			
•	l	family 2 proteins.	BL00972D 22.55 7.968e-16 210-673			
350	PR00049	WILM'S TUMOUR PROTEIN	BL00972B 9.45 1.600e-12 445-455 PR00049D 0.00 8.008e-13 121-136			
330	FR00049	SIGNATURE	PR00049D 0.00 8.0086-13 121-136 PR00049D 0.00 7.375e-12 125-140			
	ļ	SIGNATURE	PR00049D 0.00 7.3736-12 123-140			
	l		PR00049D 0.00 5.916e-11 128-143			
			PR00049D 0.00 0.7466-11 122-137			
		_	1			
			PR00049D 0.00 1.286e-10 119-134 PR00049D 0.00 8.929e-10 127-142			
	1		PR00049D 0.00 8.9296-10 127-142 PR00049D 0.00 2.678e-09 129-144			
		·	1			
	1.		PR00049D 0.00 4.051e-09 123-138 PR00049D 0.00 4.051e-09 124-139			
	j		PR00049D 0.00 4.051e-09 124-139			
350	PR00211	GLUTELIN SIGNATURE	PR00211B 0.86 7.500e-09 124-145			
350	DM00215	PROLINE-RICH PROTEIN 3.	DM00215 19.43 5.339e-10 108-141			
JJ0	DIMOUZIS	I KOLINDAIGH FROIEIN J.	DM00215 19.43 5.359e-10 106-141 DM00215 19.43 7.268e-10 112-145			
	1		DM00215 19.43 7.268e-10 112-143 DM00215 19.43 2.525e-09 106-139			
			DM00215 19.43 2.525e-09 106-139 DM00215 19.43 9.695e-09 107-140			
350	BL00048	Protamine P1 proteins.	BL00048 6.39 9.888e-09 145-172			
352	BL00518					
JJL	DTOMTO	Zinc finger, C3HC4 type (RING	BL00518 12.23 4.429e-10 214-223			

Table 3

SEQ ID Database NO: entry ID		Description	*Results
		finger), proteins.	
353	BL00518	Zinc finger, C3HC4 type (RING finger), proteins.	BL00518 12.23 4.429e-10 179-188
354	BL01009	Extracellular proteins SCP/Tpx-	BL01009D 14.19 9.341e-17 160-181
		1/Ag5/PR-1/Sc7 proteins.	BL01009A 13.75 3.769e-14 80-98
			BL01009E 13.50 5.333e-14 194-210
· · · · · · · · · · · · · · · · · · ·	<u> </u>		BL01009C 10.54 2.667e-11 127-141
354	PR00838	VENOM ALLERGEN 5 SIGNATURE	PR00838G 16.07 2.304e-14 158-178
	1.	·	PR00838D 8.73 4.452e-12 80-99 PR00838F
			10.11 7.532e-10 125-141
354	PR00837	ALLERGEN V5/TPX-1 FAMILY	PR00837C 17.21 7.429e-18 159-176
)	SIGNATURE	PR00837A 14.77 1.900e-15 80-99
			PR00837D 11.12 2.198e-13 195-209
			PR00837B 11.64 3.483e-09 127-141
356	BL00215	Mitochondrial energy transfer proteins.	BL00215A 15.82 8.500e-17 16-41
	1		BL00215B 10.44 4.900e-09 177-190
			BL00215A 15.82 6.786e-09 133-158
			BL00215B 10.44 7.300e-09 278-291
356	PR00926	MITOCHONDRIAL CARRIER	PR00926E 11.70 6.049e-13 91-110
		PROTEIN SIGNATURE	PR00926F 17.75 7.600e-11 240-263
			PR00926F 17.75 5.219e-10 18-41 PR00926D
		,	10.53 7.387e-09 246-265
357	PR00326	GTP1/OBG GTP-BINDING PROTEIN FAMILY SIGNATURE	PR00326A 8.75 7.150e-11 21-42
357	BL00113	Adenylate kinase proteins.	BL00113A 12.74 6.677e-09 22-39
357	BL01128	Shikimate kinase proteins.	BL01128A 18.84 7.802e-09 21-55
357	BL00300	SRP54-type proteins GTP-binding domain proteins.	BL00300B 20.56 1.000e-08 18-64
358	BL00972	Ubiquitin carboxyl-terminal hydrolases	BL00972A 11.93 6.318e-19 324-342
,		family 2 proteins.	BL00972D 22.55 3.903e-16 170-194
			BL00972B 9.45 1.600e-12 405-415
364	DM00215	PROLINE-RICH PROTEIN 3.	DM00215 19.43 1.482e-10 355-388
364	PR00217	43 KD POSTSYNAPTIC PROTEIN SIGNATURE	PR00217C 10.91 4.600e-10 302-318
365	BL00518	Zinc finger, C3HC4 type (RING finger), proteins.	BL00518 12.23 2.800e-11 125-134
365	BL00415	Synapsins proteins.	BL00415N 4.29 2.839e-09 387-431
365	DM00215	PROLINE-RICH PROTEIN 3.	DM00215 19.43 7.706e-11 377-410
			DM00215 19.43 8.412e-11 333-366
			DM00215 19.43 2.678e-09 356-389
			DM00215 19.43 5.138e-09 376-409
365	BL01102	Prokaryotic dksA/traR C4-type zinc finger.	BL01102 15.99 5.705e-09 109-135
365	PR00211	GLUTELIN SIGNATURE	PR00211B 0.86 5.959e-11 407-428
		•	PR00211B 0.86 2.212e-10 401-422
			PR00211B 0.86 9.500e-09 336-357
365	PR00049	WILM'S TUMOUR PROTEIN SIGNATURE	PR00049D 0.00 9.695e-09 335-350
367	PD00126	PROTEIN REPEAT DOMAIN TPR NUCLEA.	PD00126A 22.53 8.448e-09 2-23

Table 3

SEQ ID	Database	Description	*Results		
NO:	entry ID	<u> </u>			
370	BL00028	Zinc finger, C2H2 type, domain	BL00028 16.07 7.353e-14 157-174 BL00028		
		proteins.	16.07 1.000e-13 269-286 BL00028 16.07		
]		8.200e-13 493-510 BL00028 16.07 3.739e-		
			12 213-230 BL00028 16.07 6.478e-12 381-		
	l		398 BL00028 16.07 1.346e-11 185-202		
		,	BL00028 16.07 2.385e-11 129-146 BL00028		
	}		16.07 2.385e-11 325-342 BL00028 16.07		
		İ ·	5.154e-11 241-258 BL00028 16.07 9.654e-		
	•		11 437-454 BL00028 16.07 1.300e-10 297-		
			314 BL00028 16.07 9.100e-10 409-426		
L	<u> </u>	·	BL00028 16.07 9.100e-10 465-482		
370	PD00066	PROTEIN ZINC-FINGER METAL-	PD00066 13.92 2.385e-15 229-242 PD00066		
	1	BINDL	13.92 3.077e-15 145-158 PD00066 13.92		
	l		8.800e-14 173-186 PD00066 13.92 3.500e-		
	}		13 369-382 PD00066 13,92 8,500e-13 341-		
)		354 PD00066 13.92 9.133e-12 397-410		
			PD00066 13.92 2.174e-11 313-326 PD00066		
	İ		13.92 3.348e-11 453-466 PD00066 13.92		
•	1	·	3.739e-11 481-494 PD00066 13.92 7.214e-		
	ļ	`	11 257-270 PD00066 13.92 2.038e-10 425-		
			438 PD00066 13.92 6.538e-10 201-214		
	[PD00066 13.92 5.200e-09 285-298		
370	DM01970	0 kw ZK632.12 YDR313C	DM01970B 8.60 6.201e-09 265-278		
	1	ENDOSOMAL III.	·		
370	PR00048	C2H2-TYPE ZINC FINGER	PR00048A 10.52 1.474e-11 462-476		
	ļ	SIGNATURE	PR00048A 10.52 6.684e-11 182-196		
	[PR00048A 10.52 2.957e-10 434-448		
	<u>'</u>		PR00048B 6.02 5.500e-10 338-348		
•			PR00048A 10.52 6.478e-10 350-364		
			PR00048B 6.02 6.187e-10 226-236		
	.		PR00048A 10.52 6.870e-10 490-504		
•	[PR00048A 10.52 8.826e-10 406-420		
	ł ·		PR00048B 6.02 3.842e-09 170-180		
	1		PR00048B 6.02 4.316e-09 366-376		
			PR00048B 6.02 4.789e-09 478-488		
	1		PR00048B 6.02 7.632e-09 142-152		
	{		PR00048A 10.52 8.122e-09 126-140		
			PR00048B 6.02 9.053e-09 450-460		
371	BL01019	ADP-ribosylation factors family	BL01019B 19.49 6.276e-21 95-150		
		proteins.	BL01019A 13.20 8.453e-17 51-91		
371	PR00328	GTP-BINDING SAR1 PROTEIN	PR00328C 13.16 8.481e-13 78-104		
	<u> </u>	SIGNATURE	PR00328D 12.56 3.357e-11 123-145		
371	BL01115	GTP-binding nuclear protein ran proteins.	BL01115A 10.22 8.119e-11 21-65		
373	BL00028	Zinc finger, C2H2 type, domain proteins.	BL00028 16.07 4.522e-12 208-225		
373	PD00066	PROTEIN ZINC-FINGER METAL-	PD00066 13.92 7.000e-13 194-207 PD00066		
		BINDL	13.92 7.000e-13 224-237 PD00066 13.92		
	1		7.000e-12 254-267		
373	PR00048	C2H2-TYPE ZINC FINGER	PR00048A 10.52 1.391e-10 205-219		
	,	, - <u>-</u>	A A C C C C C C C C C C C C C C C C C		

Table 3

SEQ ID Databa NO: entry I		Description	*Results		
		SIGNATURE	PR00048B 6.02 6.063e-10 221-231		
374	PR00308	TYPE I ANTIFREEZE PROTEIN	PR00308A 5.90 7.288e-11 533-548		
		SIGNATURE	PR00308A 5.90 8.835e-09 534-549		
377	PD02784	PROTEIN NUCLEAR	PD02784B 26.46 7.538e-09 147-190		
	<u> </u>	RIBONUCLEOPROTEIN.			
378	PD01351	PROTEIN REPEAT	PD01351A 8.69 7.469e-09 155-166		
		NEUROFILAMENT TRIPL.	1 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2		
380	PF00094	von Willebrand factor type D domain	PF00094C 12.88 1.918e-09 43-53		
		proteins.			
380	BL01208	VWFC domain proteins.	BL01208B 15.83 3.667e-11 120-135		
			BL01208B 15.83 1.973e-09 178-193		
380	PD02138	PRECURSOR GLYCOPROTEIN	PD02138A 27.60 9.057e-09 20-69		
		SIGNAL CELL.			
381	BL01105	Ribosomal protein L35Ae proteins.	BL01105B 12.95 7.930e-13 43-83		
384	PR00049	WILM'S TUMOUR PROTEIN	PR00049D 0.00 9.205e-10 10-25 PR00049D		
		SIGNATURE	0.00 1.915e-09 9-24		
385	BL01115	GTP-binding nuclear protein ran proteins.	BL01115A 10.22 8.909e-13 34-78		
385	BL00905	GTP1/OBG family proteins.	BL00905D 15.00 5.313e-09 140-155		
385	PR00449	TRANSFORMING PROTEIN P21	PR00449C 17.27 3.209e-19 75-98		
		RAS SIGNATURE	PR00449A 13.20 1.000e-17 34-56		
			PR00449D 10.79 3.368e-13 139-153		
			PR00449B 14.34 8.364e-11 57-74 PR00449E		
	İ	,	13.50 8.286e-09 174-197		
386	BL00115	Eukaryotic RNA polymerase II	BL00115Z 3.12 7.977e-10 397-446		
	<u> </u>	heptapeptide repeat proteins.			
386	PR00041	CAMP RESPONSE ELEMENT	PR00041F 8.53 9.365e-09 256-274		
		BINDING (CREB) PROTEIN	·		
		SIGNATURE			
388	PF00646	F-box domain proteins.	PF00646A 14.37 9.036e-10 28-42		
389	BL00036	bZIP transcription factors basic domain proteins.	BL00036 9.02 6.294e-12 81-94		
389	PR00042	FOS TRANSFORMING PROTEIN	PR00042C 8.29 8.105e-13 82-99 PR00042D		
		SIGNATURE	8.97 9.895e-10 100-122		
389	BL00224	Clathrin light chain proteins.	BL00224B 16.94 3.373e-09 70-123		
389	PR00043	JUN TRANSCRIPTION FACTOR	PR00043B 8.73 9.596e-09 81-98		
	1	SIGNATURE			
390	PF00622	Domain in SPla and the RYanodine	PF00622B 21.00 2.500e-13 85-107		
		Receptor.			
391	BL00564	Argininosuccinate synthase proteins.	BL00564A 19.93 6.114e-09 7-44		
392	PR00048	C2H2-TYPE ZINC FINGER	PR00048A 10.52 7.750e-14 230-244		
200	D7 00000	SIGNATURE	PR00048A 10.52 4.316e-11 202-216		
392	BL00028	Zinc finger, C2H2 type, domain	BL00028 16.07 2.125e-15 205-222 BL00028		
		proteins.	16.07 1.391e-12 233-250 BL00028 16.07		
200	PDOCOCC	DROWNING TO TO THE TOTAL TO THE	3.400e-10 177-194		
392	PD00066	PROTEIN ZINC-FINGER METAL-	PD00066 13.92 3.000e-13 193-206 PD00066		
200	2722	BINDI.	13.92 3.423e-10 221-234		
393	PF00622	Domain in SPla and the RYanodine	PF00622B 21.00 1.391e-16 132-154		
		Receptor.			

Table 3

NO: entry ID BL00028 Zinc finger, C2H2 type, domain proteins. 16.07 2.029e-09 789-806 393 PR00048 C2H2-TYPE ZINC FINGER PR00048A 10.52 2.800e-09 758-SIGNATURE PR00501A 8.25 1.409e-09 537-55 M00099 4 kw A55R REDUCTASE TERMINAL DIHYDROPTERIDINE PR00399A 9.52 3.133e-19 146-10 PR00399 EXPRANTAGE PR00399B 14.73 4.375e-09 415 PR00399B 14.27 7.750e-16 161- PR00399B 14.27 7.750e-16 161- PR00399B 14.27 7.750e-16 161- PR00399B 14.27 7.750e-16 161- PR00399B 14.27 7.750e-16 161- PR00399B 14.27 7.750e-16 161- PR00399B 14.27 7.750e-16 161- PR00399B 14.27 7.750e-16 161- PR00399B 14.27 7.750e-16 161- PR00399B 14.27 7.750e-16 161- PR00360B 13.61 8.269e-13 201- PR00360B 13.61 8.269e-13 201- PR00360B 13.61 8.270e-10 311- PR00360B 13.61 8.270	772 51 -425 62 238 175
PR00048 C2H2-TYPE ZINC FINGER PR00048A 10.52 2.800e-09 758-SIGNATURE PR00501A 8.25 1.409e-09 537-55	772 51 -425 62 238 175
SIGNATURE PR00501 KELCH REPEAT SIGNATURE PR00501A 8.25 1.409e-09 537-53	51 -425 62 238 175
PR00501 KELCH REPEAT SIGNATURE PR00501A 8.25 1.409e-09 537-5.	425 62 238 175
DM00099	425 62 238 175
PR00399 SYNAPTOTAGMIN SIGNATURE PR00399A 9.52 3.133e-19 146-16 PR00399C 12.82 8.200e-17 222-2 PR00399B 14.27 7.750e-16 161-1 PR00399D 14.48 4.000e-14 242-2 PR00399D 14.48 4.000e-14 242-2 PR00360B 13.61 8.269e-13 201-2 PR00360B 13.61 8.269e-13 201-2 PR00360B 13.61 8.269e-13 201-2 PR00360B 13.61 5.217e-12 340-2 PR00360A 14.59 5.207e-10 311-3 PR00360A 14.59 5.207e-10 311-3 PR00360A 14.59 5.207e-10 311-3 PR00360A 14.59 5.207e-10 311-3 PR00360A 14.59 5.207e-10 311-3 PR00360B 13.61 8.214 7.231e-21 558-3 PR00168B 11.83 2.000e-09 306-3 PR00168B 11.83 2.000e-09 306-3 PR00168B 11.33 1.000e-11 185-3 PR00168B 11.33 1.000e-11 185-3 PR00168B 13.34 9.3534e-10 52-10 PR00791B 28.49 3.534e-10 52-10 PR00791B 28.49 3.534e-10 52-10 PR00791B 28.49 3.534e-10 52-10 PR00078B 13.14 4.130e-09 45-53 PR00078B 13.14 4.130e-09 45-53 PR00023B 14.20 3.077e-11 48-58 14.20 3.769e-11 176-186 PR0002 PR00023B 14.20 3.077e-11 48-58 14.20 3.769e-11 176-186 PR0002 PR00023B 14.20 3.077e-11 48-58 PR00023B 14.20 3.077e	238 175
PR00399C 12.82 8.200e-17 222-2 PR00399B 14.27 7.750e-16 161-2 PR00399D 14.48 4.000e-14 242-3 PR00399D 14.48 4.000e-14 242-3 PR00360B 13.61 8.269e-13 201-3 PR00360A 14.59 2.800e-12 174-3 PR00360B 13.61 5.217e-12 340-3 PR00360A 14.59 5.207e-10 311-3 PF00168C 27.49 5.500e-18 323-3 PF00168C 27.49 5.500e-18 323-3 PF00168B 11.83 2.000e-09 306-3 PF00168B 11.83 2.000e-09 306-3 PF00168B 11.83 2.000e-09 306-3 PF00168B 11.33 1.000e-11 185-3 PF00168B 11.33 1.000e-11 185-3 PF00791 Domain present in ZO-1 and Unc5-like netrin receptors. PF00791B 28.49 3.534e-10 52-10 PF00791B 28.49 3.534e-10 52-10 PF000791B 28.49 3.534e-10 52-10 PF000791B 28.49 3.534e-10 52-10 PF000791B 28.49 3.534e-10 52-10 PF000791B 28.49 3.534e-10 52-10 PF000791B 28.49 3.534e-10 52-10 PF000791B 28.49 3.534e-10 52-10 PF000791B 28.49 3.534e-10 52-10 PF000791B 28.49 3.534e-10 52-10 PF000791B 28.49 3.534e-10 52-10 PF000791B 28.49 3.534e-10 52-10 PF000791B 28.49 3.534e-10 52-10 PF000791B 28.49 3.534e-10 52-10 PF000791B 28.49 3.534e-10 52-10 PF000791B 28.49 3.534e-10 52-10 PF000791B 28.49 3.534e-10 52-10 PF00791B 28.49 3.534e-10 52-10 PF00791B 28.49 3.534e-10 52-10 PF00791B 28.49 3.534e-10 52-10 PF00791B 28.49 3.534e-10 52-10 PF00791B 28.49 3.534e-10 52-10 PF00791B 28.49 3.534e-10 52-10 PF00791B 28.49 3.534e-10 52-10 PF00791B 28.49 3.534e-10 52-10 PF00791B 28.49 7.291e-10 88-14 PF00791B 28.49 7.291e-10 88-14 PF00791B 28.49 7.291e-10 88-14 PF00791B 28.49 7.291e-10 88-14 PF00791B 28.49 7.291e-10 88-14 PF00791B 28.49 7.291e-10 88-14 PF00791B 28.49 7.291e-10 88-14 PF00791B 28.49 7.291e-10 88-14 PF00791B 28.49 7.291e-10 88-14 PF00791B 28.49 7.291e-10 88-14 PF00791B 28.49 7.291e-10 88-14 PF00791B 28.49 7.291e-10 88-14 PF00791B 28.49 7.291e-10 88-14 PF00791B 28.49 7.291e-10 88-14 PF00791B 28.49 7.291e-10 88-14 PF00791B 28.49 7.291e-10 88-14 PF00791B 28.49 7.291e-10 88-14 PF00791B 28.49 7.291e-10 88-14 PF00791B 28.49 7.291e-10 88-14 PF00791B 28.49	238 175
PR00399D 14.48 4.000e-14 242-395	
PR00399D 14.48 4.000e-14 242-395	
PR00360 C2 DOMAIN SIGNATURE	253
PR00360A 14.59 2.800e-12 174- PR00360B 13.61 5.217e-12 340-3 PR00360A 14.59 5.207e-10 311-3 PR00360A 14.59 5.207e-10 311-3 PR00360A 14.59 5.207e-10 311-3 PF00168C 27.49 5.500e-18 323-3 PF00168B 11.83 2.000e-09 306-3 PF00168B 11.83 2.000e-09 306-3 PF00168B 11.83 2.000e-09 306-3 PF00168B 11.83 2.000e-09 306-3 PF00168B 11.83 2.000e-09 306-3 PF00168B 11.83 2.000e-09 306-3 BL01013B 11.33 1.000e-11 185-3 PF00791 Domain present in ZO-1 and Unc5-like netrin receptors. PF00791B 28.49 3.534e-10 52-10 PF00078B 13.14 3.739e-09 78-9 PD00078B 13.14 4.130e-09 45-53 PF00023 Ank repeat proteins. PF00023B 14.20 3.077e-11 48-58 14.20 3.769e-11 176-186 PF0002 7.429e-09 85-101 PF00791B 28.49 4.455e-10 55-71 PF00791B 28.49 4.455e-11 55-11 PF00791B 28.49 7.291e-10 88-14 PF00791B 28.49 7.291e-10 88-14 PF00791B 28.49 7.291e-10 88-14 PF00791B 28.49 7.291e-10 88-14 PF00791B 28.49 7.291e-10 88-14 PF00791B 28.49 7.291e-10 88-14 PF00791B 28.49 7.291e-10 88-14 PF00791B 28.49 7.291e-10 88-14 PF00791B 28.49 7.291e-10 88-14 PF00791B 28.49 7.291e-10 88-14 PF00791B 28.49 7.291e-10 88-14 PF00791B 28.49 7.291e-10 88-14 PF00791B 28.49 7.291e-10 88-14 PF00791B 28.49 7.291e-10 88-14 PF00791B 28.49 7.291e-10 88-14 PF00791B 28.49 7.291e-10 88-14 PF00791B 28.49 7.291e-10 88-14	
PR00360B 13.61 5.217e-12 340-2 PR00360A 14.59 5.207e-10 311-3 PF00168 C2 domain proteins. PF00168C 27.49 5.500e-18 323-3 PF00168B 11.83 2.000e-09 306-3 PF00168B 11.83 2.000e-09 306-3 PF00168B 11.83 2.000e-09 306-3 PF00168B 11.83 2.000e-09 306-3 BL01013A 25.14 7.231e-21 558-3 BL01013B 11.33 1.000e-11 185-3 PF00791 Domain present in ZO-1 and Unc5-like netrin receptors. PF00791B 28.49 3.534e-10 52-10 PF00791B 28.49 3.534e-10 52-10 PF00791B 28.49 3.534e-10 52-10 PF00791B 13.14 3.739e-09 78-9 PD00078B 13.14 4.130e-09 45-53 PF00023B 14.20 3.077e-11 48-58 14.20 3.769e-11 176-186 PF0002 7.429e-09 85-101 PF00791B 28.49 4.455e-11 55-71 PF00791B 28.49 4.455e-11 55-71 PF00791B 28.49 7.291e-10 88-14 PF00791B 28.49 7.291e-10 88-14 PR00450 RECOVERIN FAMILY SIGNATURE PR00450D 16.58 8.986e-11 161-1	
PR00360A 14.59 5.207e-10 311-395 PF00168	
395 PF00168 C2 domain proteins. PF00168C 27.49 5.500e-18 323-3 PF00168B 11.83 2.000e-09 306-3 PF00168B 11.83 2.000e-09 306-3 PF00168B 11.83 2.000e-09 306-3 PF00168B 11.83 2.000e-09 306-3 PF00168B 11.83 2.000e-09 306-3 PF00168B 11.83 2.000e-09 306-3 PF00091 PF0	
PF00168B 11.83 2.000e-09 306-3 PF00168B 11.83 2.000e-09 306-3 PF00791	
BL01013 Oxysterol-binding protein family proteins. BL01013A 25.14 7.231e-21 558-101	17
396 PF00791 Domain present in ZO-1 and Unc5-like netrin receptors. PF00791B 28.49 3.534e-10 52-10 string receptors. 396 PD00078 REPEAT PROTEIN ANK NUCLEAR ANK YR. PD00078B 13.14 9.000e-11 173-PD00078B 13.14 3.739e-09 78-9 PD00078B 13.14 4.130e-09 45-53 PD00078B 13.14 4.130e-09 45-53 PD00078B 13.14 4.130e-09 45-53 PD00078B 13.14 4.130e-09 45-53 PD00078B 13.14 4.130e-09 45-53 PF00023B 14.20 3.077e-11 48-58 PF00023B 14.20 3.0769e-11 176-186 PF0002 TA.429e-09 85-101 397 PF00023 Ank repeat proteins. PF00023A 16.03 1.750e-10 55-71 PF00791B 28.49 4.455e-11 55-11 netrin receptors. 398 BL00422 Granins proteins. BL00422C 16.18 5.787e-10 134-400 PR00450 400 PR00450 RECOVERIN FAMILY SIGNATURE PR00450D 16.58 8.986e-11 161-10	
396 PF00791 Domain present in ZO-1 and Unc5-like netrin receptors. PF00791B 28.49 3.534e-10 52-10 section receptors. 396 PD00078 REPEAT PROTEIN ANK NUCLEAR ANK YR. PD00078B 13.14 9.000e-11 173-PD00078B 13.14 4.130e-09 78-9.PD00078B 13.	196
396 PD00078 REPEAT PROTEIN ANK NUCLEAR ANK YR. PD00078B 13.14 9.000e-11 173-174 173-	
PD00078B 13.14 4.130e-09 45-58 PF00023	186
396 PF00023 Ank repeat proteins. PF00023B 14.20 3.077e-11 48-58 14.20 3.769e-11 176-186 PF0002 7.429e-09 85-101 397 PF00023 Ank repeat proteins. PF00023A 16.03 1.750e-10 55-71 PF00023A 16.03 1.750e-10 55-71 PF00023A 16.03 1.750e-10 55-71 PF00023A 16.03 1.750e-10 55-71 PF00023A 16.03 1.750e-10 55-71 PF00023A 16.03 1.750e-10 55-71 PF000023A 16.03 1.750e-10 55-71 PF000023A 16.03 1.750e-10 10 55-71 PF000023A 16.03 1.750e-10 10 55-71 PF000023A 16.03 1.750e-10 10 55-71 PF000023A 16.03 1.750e-10 10 55-71 PF000023A 16.03 1.750e-10 10 55-71 PF000023A 16.03 1.750e-10 10 55-71 PF000023A 16.03 1.750e-10 10 55-71 PF000023A 16.03 1.750e-10 10 55-71 PF000023A 16.03 1.750e-10 10 55-71 PF000023A 16.03 1.750e-10 10 55-71 PF000023A 16.03 1.750e-10 10 55-71 PF000023A 16.03 1.750e-10 10 55-71 PF000023A 16.03 1.750e-10 10 55-71 PF000023A 16.03 1.750e-10 10 55-71 PF000023A 16.03 1.750e-10 10 55-71 PF000023A 16.03 1.750e-10 10 55-71 PF000000000000000000000000000000000000	1
14.20 3.769e-11 176-186 PF0002 7.429e-09 85-101 397 PF00023 Ank repeat proteins. PF00023A 16.03 1.750e-10 55-71 397 PF00791 Domain present in ZO-1 and Unc5-like PF00791B 28.49 4.455e-11 55-11 netrin receptors. PF00791B 28.49 7.291e-10 88-14 398 BL00422 Granins proteins. BL00422C 16.18 5.787e-10 134-1400 PR00450 RECOVERIN FAMILY SIGNATURE PR00450D 16.58 8.986e-11 161-140 PR00450D 16.58 PR00	3
7.429e-09 85-101	PF00023B
397 PF00023 Ank repeat proteins. PF00023A 16.03 1.750e-10 55-71 397 PF00791 Domain present in ZO-1 and Unc5-like netrin receptors. PF00791B 28.49 4.455e-11 55-11 398 BL00422 Granins proteins. BL00422C 16.18 5.787e-10 134-14 400 PR00450 RECOVERIN FAMILY SIGNATURE PR00450D 16.58 8.986e-11 161-1	3A 16.03
397 PF00791 Domain present in ZO-1 and Unc5-like netrin receptors. PF00791B 28.49 4.455e-11 55-11 PF00791B 28.49 7.291e-10 88-14 398 BL00422 Granins proteins. BL00422C 16.18 5.787e-10 134-14 400 PR00450 RECOVERIN FAMILY SIGNATURE PR00450D 16.58 8.986e-11 161-1	
netrin receptors. PF00791B 28.49 7.291e-10 88-14 398 BL00422 Granins proteins. BL00422C 16.18 5.787e-10 134-14 400 PR00450 RECOVERIN FAMILY SIGNATURE PR00450D 16.58 8.986e-11 161-14	
398 BL00422 Granins proteins. BL00422C 16.18 5.787e-10 134-1 400 PR00450 RECOVERIN FAMILY SIGNATURE PR00450D 16.58 8.986e-11 161-1	.0
400 PR00450 RECOVERIN FAMILY SIGNATURE PR00450D 16.58 8.986e-11 161-	3
	62
400 BI 00470 Bhoshol agtom / discrelely regard his discretely 12 57 4 272 15 207	181
400 BL00479 Phorbol esters / diacylglycerol binding BL00479B 12.57 4.273e-15 287-3	303
domain proteins. BL00479A 19.86 2.667e-14 261-2	284
BL00479B 12.57 1.360e-10 351-3	367
400 PR00171 CLASS III CYTOCHROME C PR00171D 7.30 9.419e-10 334-34 SIGNATURE	1 2
400 BL00018 EF-hand calcium-binding domain BL00018 7.41 3.348e-09 223-236 proteins.	
400 PF00781 Diacylglycerol kinase catalytic domain PF00781F 16.43 1.000e-40 600-1	~~
proteins (presumed). PF00781B 12.07 8.364e-35 454-4	yy
PF00781D 11.11 3.077e-30 532-1	
PF00781C 9.69 5.034e-19 506-52	186
PF00781E 12.45 2.385e-17 124-5	186 118
PF00781G 10.09 6.211e-17 678-0	186 118 21
PF00781H 12.20 1.750e-16 770-7	866 118 21 83
PF00781A 6.42 3.667e-09 354-36	186 118 21 83 592
401 PR00049 WILM'S TUMOUR PROTEIN PR00049D 0.00 7.407e-09 325-34 SIGNATURE	886 118 21 883 592 782
402 DM01117 2 kw TRANSPOSASE WITHIN DM01117A 11.17 7.750e-09 364	186 118 21 83 592 782

Table 3

SEQ ID NO:	Database entry ID	Description	*Results		
		TRANSPOSITION VASOTOCIN.			
403	DM01206	CORONAVIRUS NUCLEOCAPSID	DM01206B 10.69 9.286e-12 724-744		
	•	PROTEIN.	DM01206B 10.69 3.466e-10 726-746		
]		DM01206B 10.69 9.630e-10 722-742		
İ			DM01206B 10.69 7.152e-09 718-738		
	<u></u>		DM01206B 10.69 8.861e-09 728-748		
403	BL00048	Protamine P1 proteins.	BL00048 6.39 4.197e-10 722-749 BL00048		
]		6.39 5.500e-10 731-758 BL00048 6.39		
•	1.		6.329e-10 729-756 BL00048 6.39 9.171e-10		
1			730-757 BL00048 6.39 4.038e-09 728-755		
			BL00048 6.39 8.538e-09 724-751 BL00048		
			6.39 9.438e-09 716-743		
403	.PD00289	PROTEIN SH3 DOMAIN REPEAT PRESYNA.	PD00289 9.97 9.690e-09 130-144		
404	PD01066	PROTEIN ZINC FINGER ZINC-	PD01066 19.43 1.353e-27 31-70		
		FINGER METAL-BINDING NU.			
404	PD00066	PROTEIN ZINC-FINGER METAL-	PD00066 13.92 5.154e-15 274-287 PD00066		
	1	BINDI.	13.92 7.600e-14 246-259 PD00066 13.92		
	1		8.200e-14 302-315 PD00066 13.92 3.143e-		
			12 218-231 PD00066 13.92 4.000e-12 190-		
			203 PD00066 13.92 2.800e-09 330-343		
404	BL00028	Zinc finger, C2H2 type, domain	BL00028 16.07 7.261e-12 230-247 BL00028		
		proteins.	16.07 9.171e-12 342-359 BL00028 16.07		
]		4.300e-10 314-331 BL00028 16.07 7.000e-		
			10 174-191 BL00028 16.07 3.314e-09 202-		
			219 BL00028 16.07 6.400e-09 286-303		
404	PR00048	C2H2-TYPE ZINC FINGER	PR00048A 10.52 4.214e-13 339-353		
		SIGNATURE	PR00048A 10.52-3.209e-12 227-241		
			PR00048A 10.52 1.947e-11 311-325		
			PR00048A 10.52 4.522e-10 171-185		
			PR00048B 6.02 2.895e-09 299-309		
]		PR00048A 10.52 4.600e-09 199-213		
			PR00048B 6.02 1.000e-08 187-197		
			PR00048B 6.02 1.000e-08 271-281		
406	BL00610	Sodium:neurotransmitter symporter	BL00610A 17.73 1.000e-40 68-118		
		family proteins.	BL00610B 23.65 1.000e-40 132-182		
			BL00610C 12.94 1.000e-40 225-277		
			BL00610D 20.97 1.000e-40 291-344		
			BL00610F 29.02 6.143e-36 540-157		
	·		BL00610E 20.34 3.209e-35 448-491		
10.5			BL00610G 12.89 2.200e-15 173-196		
406	PR00176	SODIUM/NEUROTRANSMITTER	PR00176C 10.84 6.226e-23 141-168		
		SYMPORTER SIGNATURE	PR00176A 16.82 1.450e-22 68-90 PR00176F		
			10.73 8.667e-20 452-472 PR00176B 7.31		
			7.000e-18 97-117 PR00176D 9.02 1.000e-17		
			252-270 PR00176E 11.41 2.756e-15 334-355		
	ļ		PR00176H 15.27 7.353e-15 131-590		
			PR00176G 12.48 5.615e-14 529-112		
407	DM00179	w KINASE ALPHA ADHESION T- CELL.	DM00179 13.97 5.304e-09 111-121		

Table 3

SEQ ID	Database	Description	*Results			
NO:	entry ID					
408	PR00187	DNAJ PROTEIN FAMILY	PR00187B 13.48 1.800e-16 45-66			
	<u> </u>	SIGNATURE	PR00187A 12.84 6.700e-12 15-35			
408	BL00198	Nt-dnaJ domain proteins.	BL00198B 15.11 9.217e-15 45-66			
	<u> </u>		BL00198A 8.07 2.459e-11 19-36			
409	PR00927	ADENINE NUCLEOTIDE	PR00927E 14.93 4.136e-11 246-268			
<u></u>	<u> </u>	TRANSLOCATOR 1 SIGNATURE				
409	BL00215	Mitochondrial energy transfer proteins.	BL00215A 15.82 9.735e-14 11-36			
	ļ		BL00215A 15.82 5.787e-11 108-133			
			BL00215B 10.44 6.211e-11 258-271			
			BL00215A 15.82 5.018e-09 211-236			
409	PR00926	MITOCHONDRIAL CARRIER PROTEIN SIGNATURE	PR00926D 10.53 5.355e-09 19-38			
410	PD00066	PROTEIN ZINC-FINGER METAL-	PD00066 13.92 6.400e-17 411-424 PD00066			
		BINDL	13.92 8.200e-17 327-340 PD00066 13.92			
			5.154e-15 271-284 PD00066 13.92 2.800e-			
			14 215-228 PD00066 13.92 9.000e-13 355-			
	· ·	·	368 PD00066 13.92 6.143e-12 439-452			
			PD00066 13.92 6.478e-11 187-200 PD00066			
410	DT 00000	7: 6	13.92 9.217e-11 243-256			
410	BL00028	Zinc finger, C2H2 type, domain	BL00028 16.07 2.588e-14 227-244 BL00028			
		proteins.	16.07 6.824e-14 395-412 BL00028 16.07			
			7.882e-14 171-188 BL00028 16.07 2.350e-			
			13 339-356 BL00028 16.07 7.300e-13 283-			
		_	300 BL00028 16.07 7.300e-13 367-384			
			BL00028 16.07 2.565e-12 423-440 BL00028 16.07 7.261e-12 199-216 BL00028 16.07			
			7.261e-12 311-328 BL00028 16.07 8.435e-			
			12 451-468 BL00028 16.07 2.038e-11 255-			
			272 BL00028 16.07 9.400e-10 143-160			
410	PR00048	C2H2-TYPE ZINC FINGER	PR00048A 10.52 3.250e-14 280-294			
, ,	1	SIGNATURE	PR00048A 10.52 8.500e-14 336-350			
			PR00048A 10.52 7.429e-13 252-266			
			PR00048A 10.52 8.714e-13 448-462			
			PR00048A 10.52 9.357e-13 392-406			
			PR00048A 10.52 1.000e-12 168-182			
			PR00048A 10.52 2.059e-12 420-434			
			PR00048B 6.02 8.615e-11 408-418			
			PR00048B 6.02 7.188e-10 268-278			
•	·		PR00048B 6.02 7.188e-10 380-390			
			PR00048B 6.02 9.438e-10 296-306			
			PR00048B 6.02 1.000e-09 324-334			
		'	PR00048B 6.02 1.474e-09 352-362			
			PR00048B 6.02 3.842e-09 212-222			
			PR00048B 6.02 5.263e-09 436-446			
411	BL00018	EF-hand calcium-binding domain proteins.	BL00018 7.41 5.500e-10 63-76			
413	PR00014	FIBRONECTIN TYPE III REPEAT SIGNATURE	PR00014C 15.44 4.600e-10 73-92			
414	PR00806	VINCULIN SIGNATURE	PR00806A 6.63 1.493e-09 785-796			
414	PR00048	C2H2-TYPE ZINC FINGER	PR00048A 10.52 4.240e-09 41-55			

Table 3

SEQ ID NO:	Database entry ID	Description	*Results		
	T	SIGNATURE			
414	PR00049	WILM'S TUMOUR PROTEIN	PR00049D 0.00 9.546e-11 781-796		
		SIGNATURE	PR00049D 0.00 1.205e-10 263-278		
			PR00049D 0.00 4.356e-09 785-800		
414	BL00412	Neuromodulin (GAP-43) proteins.	BL00412D 16.54 4.673e-09 420-471		
414	BL00422	Granins proteins.	BL00422C 16.18 6.318e-11 439-467		
	1	•	BL00422C 16.18 9.809e-10 440-468		
	l	•	BL00422C 16.18 6.294e-09 441-469		
			BL00422C 16.18 6.209e-09 438-466		
414	PR00910	LUTEOVIRUS ORF6 PROTEIN SIGNATURE	PR00910A 2.51 8.179e-09 265-278		
414	DM00215	PROLINE-RICH PROTEIN 3.	DM00215 19.43 4.203e-09 770-803		
747	211100213	I KOMIN MOIT I KOTEM J.	DM00215 19.43 4.2056-09 770-805 DM00215 19.43 9.0856-09 245-278		
414	BL00028	Zinc finger, C2H2 type, domain	BL00028 16.07 1.257e-09 44-61 BL00028		
747	D100020	proteins.	16.07 2.543e-09 175-192 BL00028 16.07		
	1	browns.	6.143e-09 119-136 BL00028 16.07 9.743e-		
	ŀ		09 147-164		
415	PF00622	Receptor.Domain in SPla and the	PF00622B 21.00 1.000e-13 331-353		
.10	1100022	RYanodine	PF00622C		
415	BL00518	Zinc finger, C3HC4 type (RING	BL00518 12.23 3.400e-11 31-40		
	2200510	finger), proteins.	55.00510 12.25 5.4000-11 51-40		
416	PF00780	Domain found in NIK1-like kinases,	PF00780B 23.03 5.929e-33 442-485		
	1	mouse citron and yeast ROM.			
416	PR00109	TYROSINE KINASE CATALYTIC	PR00109B 12.27 5.235e-12 211-230		
		DOMAIN SIGNATURE			
416	BL00107	Protein kinases ATP-binding region	BL00107A 18.39 5.200e-22 211-242		
	1	proteins.	BL00107B 13.31 9.308e-12 283-299		
416	BL00239	Receptor tyrosine kinase class II proteins.	BL00239B 25.15 5.164e-10 145-193		
416	BL00915	Phosphatidylinositol 3- and 4-kinases	BL00915C 22.43 9.357e-10 203-242 *		
		proteins.			
417	BL00021	Kringle domain proteins.	BL00021B 13.33 1.482e-14 41-59		
		, G	BL00021D 24.56 2.122e-12 193-235		
417	PR00722	CHYMOTRYPSIN SERINE	PR00722A 12.27 7.517e-14 42-58		
		PROTEASE FAMILY (S1)	PR00722B 12.51 3.143e-10 97-112		
		SIGNATURE			
417	BL00134	Serine proteases, trypsin family,	BL00134A 11.96 6.464e-16 41-58		
-		histidine proteins.	BL00134C 13.45 2.059e-09 221-235		
417	BL00495	Apple domain proteins.	BL00495O 13.75 2.440e-09 212-241		
417	BL00672	Serine proteases, V8 family, histidine	BL00672A 9.79 9.520e-09 41-57		
. = •		proteins.			
417	PR00839	V8 SERINE PROTEASE FAMILY	PR00839B 11.20 9.753e-09 41-59		
		SIGNATURE			
418	BL01207	Glypicans proteins.	BL01207B 23.69 9.122e-28 191-237		
		• • • • • • • • • • • • • • • • • • • •	BL01207A 12.21 1.000e-16 62-78		
423	PD02870	RECEPTOR INTERLEUKIN-1	PD02870D 15.74 4.351e-09 693-728		
-		PRECURSOR.			
423	DM00179	w KINASE ALPHA ADHESION T-	DM00179 13.97 5.696e-09 793-803		
		CELL.			

Table 3

SEQ ID NO:	Database entry ID	Description	*Results			
424	BL00203	Vertebrate metallothioneins proteins.	BL00203 13.94 5.041e-09 13-59			
425	BL00107	Protein kinases ATP-binding region proteins.	BL00107A 18.39 8.141e-18 217-248			
425	BL00240	Receptor tyrosine kinase class III proteins.	BL00240E 11.56 6.040e-10 203-241			
425	PR00109	TYROSINE KINASE CATALYTIC DOMAIN SIGNATURE	PR00109B 12.27 5.814e-14 217-236 PR00109A 15.00 1.730e-09 182-196			
428	PR00141	PROTEASOME COMPONENT SIGNATURE	PR00141C 11.15 6.333e-12 234-246 PR00141D 12.45 8.615e-12 259-271 PR00141B 11.15 9.561e-12 223-235 PR00141A 11.36 2.050e-11 102-118			
428	BL00854	Proteasome B-type subunits proteins.	BL00854A 33.93 1.383e-19 99-145 BL00854C 29.92 5.235e-14 206-235 BL00854D 13.76 2.800e-09 257-267			
429	PR00245	OLFACTORY RECEPTOR SIGNATURE	PR00245A 18.03 9.413e-17 59-81 PR00245C 7.84 7.500e-16 238-254 PR00245E 12.40 2.500e-12 291-306 PR00245B 10.38 9.112e-11 177-192			
429	PR00237	RHODOPSIN-LIKE GPCR SUPERFAMILY SIGNATURE	PR00237E 13.03 7.120e-12 199-223 PR00237C 15.69 1.225e-09 104-127			
429	BL00237	G-protein coupled receptors proteins.	BL00237A 27.68 9.727e-14 90-130 BL00237D 11.23 1.273e-09 282-299			
429	PR00534	MELANOCORTIN RECEPTOR FAMILY SIGNATURE	PR00534A 11.49 6.400e-09 51-64			
430	PF00651	BTB (also known as BR-C/Ttk) domain proteins.	PF00651 15.00 1.000e-11 87-100			
430	BL00028	Zinc finger, C2H2 type, domain proteins.	BL00028 16.07 4.706e-14 474-491 BL00028 16.07 1.771e-09 502-519			
430	PD00066	PROTEIN ZINC-FINGER METAL- BINDL	PD00066 13.92 4.300e-09 490-503			
430	PR00048	C2H2-TYPE ZINC FINGER SIGNATURE	PR00048A 10.52 4.600e-09 499-513			
433	BL00086	Cytochrome P450 cysteine heme-iron ligand proteins.	BL00086 20.87 3.209e-23 430-462			
433	PR00465	E-CLASS P450 GROUP IV SIGNATURE	PR00465F 13.37 1.360e-11 400-419			
433	PR00359	B-CLASS P450 SIGNATURE	PR00359G 11.22 8.071e-10 401-417 PR00359F 24.20 2.180e-09 373-401			
433	PR00385	P450 SUPERFAMILY SIGNATURE	PR00385E 12.66 8.800e-11 440-452 PR00385D 13.11 4.429e-10 431-441 PR00385A 14.97 5.865e-09 302-320			
433	PR00464	E-CLASS P450 GROUP II SIGNATURE	PR00464G 12.41 9.000e-10 405-421 PR00464D 17.40 1.191e-09 320-338 PR00464E 18.28 6.946e-09 349-370 PR00464H 13.32 7.750e-09 427-441 PR00464C 18.84 9.014e-09 291-320 PR00464I 14.64 9.481e-09 440-464			
434	BL00216	Sugar transport proteins.	BL00216B 27.64 7.943e-19 101-151			
434	PR00171	SUGAR TRANSPORTER SIGNATURE	PR00171D 12.76 3.593e-11 413-435			

Table 3

SEQ ID NO:	Database entry ID	Description	*Results
435	PR00049	WILM'S TUMOUR PROTEIN SIGNATURE	PR00049D 0.00 8.429e-10 10-25
435	BL00028	Zinc finger, C2H2 type, domain proteins.	BL00028 16.07 4.150e-13 138-593 BL00028 16.07 6.850e-13 1010-1027 BL00028 16.07 6.087e-12 982-999 BL00028 16.07 8.615e-11 846-863 BL00028 16.07 3.100e-10 317-334 BL00028 16.07 7.000e-10 170-187 BL00028 16.07 8.500e-10 289-306 BL00028 16.07 8.800e-10 548-565
435	PD00066	PROTEIN ZINC-FINGER METAL- BINDL	PD00066 13.92 7.600e-14 998-1011 PD00066 13.92 1.000e-11 305-318 PD00066 13.92 8.826e-11 564-577 PD00066 13.92 3.400e-09 862-875
435	PR00456	RIBOSOMAL PROTEIN P2 SIGNATURE	PR00456E 3.06 5.329e-09 177-192 PR00456E 3.06 5.899e-09 140-155
435	BL00999	Streptomyces subtilisin-type inhibitors proteins.	BL00999A 14.95 7.223e-09 461-499
435	PR00048	C2H2-TYPE ZINC FINGER SIGNATURE	PR00048A 10.52 9.357e-13 573-587 PR00048A 10.52 2.421e-11 1007-1021 PR00048B 6.02 2.125e-10 561-133 PR00048A 10.52 8.043e-10 314-328 PR00048B 6.02 1.000e-09 995-1005 PR00048B 6.02 6.684e-09 302-312 PR00048A 10.52 9.280e-09 167-181
436	PR00245	OLFACTORY RECEPTOR SIGNATURE	PR00245A 18.03 2.667e-23 100-122 PR00245C 7.84 1.783e-14 232-248 PR00245D 10.47 7.070e-10 268-280
436	PR00237	RHODOPSIN-LIKE GPCR SUPERFAMILY SIGNATURE	PR00237C 15.69 8.500e-11 145-168 PR00237G 19.63 6.023e-09 266-293
436	BL00237	G-protein coupled receptors proteins.	BL00237A 27.68 2.161e-15 131-171 BL00237D 11.23 8.091e-09 276-293
437	PR00262	IL1/HBGF FAMILY SIGNATURE	PR00262A 28.26 1.000e-08 80-108
438	BL00884	Osteopontin proteins.	BL00884B 12.47 1.000e-40 50-94 BL00884C 22.45 6.187e-39 131-173 BL00884A 11.35 5.846e-32 1-31 BL00884E 11.04 8.364e-23 273-295 BL00884D 8.79 3.323e-18 255-272
438	PR00216	OSTEOPONTIN SIGNATURE	PR00216B 7.89 4.553e-34 37-67 PR00216A 10.94 8.054e-33 2-32 PR00216C 9.63 2.565e-32 67-93 PR00216G 12.39 8.676e-27 238-264 PR00216H 7.41 5.295e-22 273-293 PR00216F 11.79 3.133e-21 164-183 PR00216D 2.74 5.800e-18 104-119 PR00216E 8.44 4.405e-16 132-147

* Results include in order: Accession No., subtype, e-value, and amino acid position of the signature in the corresponding polypeptide

Table 4

SEQ ID	Pfam Model	Description	E-value	Score	No: of Pfam Domains	Position of the Domain
1	zf-CCCH	Zinc finger C-x8-C-x5-C-x3- H type	1.8e-05	31.6	1	412-438
1	zf-C3HC4	Zinc finger, C3HC4 type (RING finger)	2e-05	21.8	1	14-52
3	EMP24_GP25L	emp24/gp25L/p24 family	4.1e-105	362.6	1	22-235
6	WW	WW domain	1,2e-05	32.2	1	45-75
7	ww	WW domain	1.2e-05	32.2	1	45-75
8	Aa_trans	Transmembrane amino acid transporter protein	9.6e-64	225.2	1	71-451
9	Fe-ADH	Iron-containing alcohol dehydrogenase	9.9e-35	124.5	. 2	4-205:228- 255
10	Fe-ADH	Iron-containing alcohol dehydrogenase	9.9e-35	124.5	2	52-253:276- 303
11	Bcl-2	Apoptosis regulator proteins, Bcl-2 family	0.016	-2.1	1	257-356
12	spectrin	Spectrin repeat	1.3e-10	43.6	3	11-87:90- 197:200-291
13	Ribosomal_L18ae	Ribosomal L18ae protein family	1.9e-128	440.1	1	6-176
14	Ribosomal_L31e	Ribosomal protein L31e	2.4e-47	170.7	1	72-166
15	zf-CCCH	Zinc finger C-x8-C-x5-C-x3- H type	7.8e-16	66.0	3	342-367:371- 396:398-420
16	zf-MYND	MYND finger	1.4e-13	58.5	1	52-90
17	Sterile	Male sterility protein	1.1e-51	185.1	1	254-446
18	MgtE	Divalent cation transporter	8.6e-39	142.3	2	138-274:352- 499
19	Rap_GAP	Rap/ran-GAP	2e-124	426.7	1	400-588
19	PDZ	PDZ domain (Also known as DHR or GLGF)	2.4e-06	34.5	1	726-800
20	Rap_GAP	Rap/ran-GAP	2e-124	426.7	1	400-588
20	PDZ	PDZ domain (Also known as DHR or GLGF)	2.4e-06	34.5	1	726-800
22	SCAN	SCAN domain	1.5e-23	91.7	1	165-238
23	RhoGAP	RhoGAP domain	3e-58	206.9	1	497-649
23	FCH	Fes/CIP4 homology domain	1.2e-18	75.4	1	22-121
23	SH3	SH3 domain	2.6e-11	51.0	1	723-777
24 25	adh_zinc UDPGT	Zinc-binding dehydrogenases	1.5e-05	-25.4	1	20-336
		UDP-glucoronosyl and UDP- glucosyl transferas	1.6e-84	294.3	1	26-467
28	Ribosomal L6e	Ribosomal protein L6e	4.3e-77	269.5	1	109-239
29	Ribosomal_L11	Ribosomal protein L11	4.9e-64	226.2	1	13-144
30	tRNA-synt_le	tRNA synthetases class I (C)	1.6e-137	470.2	1	64-538
32	zf-C3HC4	Zinc finger, C3HC4 type (RING finger)	0.00041	17.6	2	33-72:165- 185
34	ras	Ras family	1.4e-77	271.2	1	35-235
34	arf	ADP-ribosylation factor family	9.3e-05	-56.3	1	17-198
36	SET	SET domain	3.2e-05	10.0	1	209-342
36	MORN	MORN repeat	0.006	23.2	3	36-58:59- 81:106-128

Table 4

SEQ ID	Pfam Model	Description	E-value	Score	No: of Pfam Domains	Position of the Domain
37	laminin_G	Laminin G domain	1.5e-11	44.7	1	55-174
37	EGF	EGF-like domain	0.0033	24.1	1	202-234
38	Sema	Sema domain	1.7e-127	436.9	1	56-489
38	Plexin_repeat	Plexin repeat	1e-06	35.7	1	507-563
38	ig	Immunoglobulin domain	0.0023	15.9	1	582-639
38	integrin_B	Integrins, beta chain	0.084	6.1	1	513-527
40	filament	Intermediate filament protein	1.6e-138	473.6	1	129-442
41	Keratin_B2	Keratin, high sulfur B2 protein	1.8e-18	74.8	2	2-138:139- 240
44	sushi	Sushi domain (SCR repeat)	3.8e-06	33.9	4	1396- 1459:1464- 1521:1525- 1590:1595- 1646
45	profilin	Profilin	4.1e-13	51.7	1	10-124
47	ubiquitin	Ubiquitin family	0.00033	20.5	1	31-99
48	BTB	BTB/POZ domain	2.6e-21	84.2	1	80-196
48	Kelch	Kelch motif	2.6e-20	80.9	4	336-382:384- 430:432- 478:582-635
48	SCP	SCP-like extracellular protein	0.015	13.0	1	1-35
49	serpin	Serpin (serine protease inhibitor)	2.4e-178	605.4	1	59-432
50	T-box	T-box	3.6e-125	429.2	1	140-331
52	7tm_1	7 transmembrane receptor (rhodopsin family)	1.2e-17	58.3	2	132-228:337- 344
53	CSD	'Cold-shock' DNA-binding domain	1.8e-16	63.6	1	42-112
53	zf-CCHC	Zinc knuckle	0.00012	28.8	2	137-154:159- 176
54	ig	Immunoglobulin domain	2.5e-07	28.7	1	34-109
55	Rap_GAP	Rap/ran-GAP	5e-18	73.3	1	287-466
57	G-gamma	GGL domain	1.8e-11	39.4	2	49-70:109-
58	T-box	T-box	8.9e-114	391.4	1	101-302
59	Gag_p10	Retroviral GAG p10 protein	9.2e-06	23.7	1	82-171
61	60s_ribosomal	60s Acidic ribosomal protein	0.0089	12.0	1	1-22
62	UPAR_LY6	u-PAR/Ly-6 domain	5.4e-05	22.3	1	8-51
63	Ribosomal_L30	Ribosomal protein L30p/L7e	0.00042	18.5	1	65-93
64	filament	Intermediate filament protein	1.1e-78	274.8	2	161-338:339- 426
65	Ribosomal_S6	Ribosomal protein S6	0.00082	7.5	1	2-96
66	PDZ	PDZ domain (Also known as DHR or GLGF)	5.1e-09	43.4	1	158-250
67	zf-C3HC4	Zinc finger, C3HC4 type (RING finger)	0.005	14.0	1	92-118
68	G-patch	G-patch domain	6.8e-07	36.3	1	26-70
69	Keratin_B2	Keratin, high sulfur B2 protein	0.037	-45.9	1	10-155
83	ig	Immunoglobulin domain	8.5e-09	33.4	2	34-89:119- 187
86	zf-C2H2	Zinc finger, C2H2 type	2.2e-71	250.6	17	182-204:210-

Table 4

SEQ ID	Pfam Model	Description	E-value	Score	No: of Pfam Domains	Position of the Domain
						232:237- 260:265- 288:315- 337:343- 365:369-
					}	392:653- 675:681- 704:709-
						733:741- 764:791- 814:820-
	,					842:848- 870:877- 899:905- 928:952-975
87	ig	Immunoglobulin domain	2.7e-35	118.7	6	36-121:162- 249:292- 375:422- 517:564- 657:704-795
88	MAP1_LC3	Microtubule associated protein 1A/1B, light	9.4e-79	275.0	1	118-221
89	WD40	WD domain, G-beta repeat	1.6e-12	55.1	4	173-215:221- 263:269- 305:1103- 1140
90	FKBP	FKBP-type peptidyl-prolyl cis-trans isomeras	1.2e-59	198.9	1	66-160
92	RPEL	RPEL repeat	6.5e-18	73.0	2	513-538:551- 576
93	transket_pyr	Transketolase, pyridine binding domain	4.6e-65	229.6	1	568-773
93	E1_dehydrog	Dehydrogenase E1 component	8.7e-23	89.1	1	193-504
95	zf-C3HC4	Zinc finger, C3HC4 type (RING finger)	8.7e-09	32.7	1	595-635
97	ig	Immunoglobulin domain	1.8e-20	71.0	3	31-88:127- 185:222-278
98	ig	Immunoglobulin domain	1.8e-20	71.0	3	24-81:120- 178:215-271
99	Patched	Patched family	6.2e-06	-369.1	1	66-935
102	zf-C2H2	Zinc finger, C2H2 type	2.3e-94	326.9	12	209-231:237- 259:265- 287:293- 315:321- 343:349- 371:377- 399:405-
		·				427:433- 455:461- 483:489- 511:594-616

Table 4

SEQ ID	Pfam Model	Description	E-value	Score	No: of Pfam Domains	Position of the Domain
102	KRAB	KRAB box	3.7e-37	136.9	1	15-77
103	zf-C2H2	Zinc finger, C2H2 type	1.2e-55	198.2	9	172-195:271- 293:299- 321:327- 349:355- 377:383- 405:411-
						433:439-
103	KRAB	KRAB box	3e-46	167.1	1	461:467-489 8-70
107	zf-CCHC	Zinc knuckle	2.4e-16	67.8	3	913-
						930:1293- 1310:1358- 1375
107	NTP_transf_2	Nucleotidyltransferase domain	4.4e-11	50.3	1	972-1065
108	zf-C2H2	Zinc finger, C2H2 type	1.6e-42	154.7	5	283:289- 311:317- 339:345- 367:373-395
109	myosin_head	Myosin head (motor domain)	0	1267.5	1	26-697
109	IQ	IQ calmodulin-binding motif	1.2e-17	72.1	4	714-734:737- 757:760- 780:789-809
110	pkinase	Protein kinase domain	1.2e-96	334.5	1	20-271
111	WD40	WD domain, G-beta repeat	1.8e-49	177.8	8	161-197:218- 253:258- 294:300- 335:341- 377:383- 428:434- 470:476-511
112	SNF2_N	SNF2 and others N-terminal domain	4.2e-78	272.9	1	1-264
112	helicase_C	Helicase conserved C- terminal domain	1.2e-24	95.4	1	326-410
113	DUF15	Domain of unknown function DUF15	0.00064	-60.4	1	132-384
114	DSPc	Dual specificity phosphatase, catalytic	0.0004	-2.9	1	141-295
114	Y_phosphatase	Protein-tyrosine phosphatase	0.0037	-26.9	1	128-295
115	Ulp1_C	Ulp1 protease family, C- terminal catalytic d	2.8e-52	187.1	1	394-587
117	Rhodanese	Rhodanese-like domain	1e-05	32.4	1	160-260
119	ABC1	ABC1 family	1.7e-40	147.9	1	318-434
122	proteasome	Proteasome A-type and B- type	7.4e-43	155.8	1	39-146
124	Ribosomal_L9	Ribosomal protein L9	3.1e-05	-3.4	1	94-240
125	RIO1	RIO1/ZK632.3/MJ0444 family	7.8e-80	278.6	1	193-387
128	abhydrolase	alpha/beta hydrolase fold	4.5e-20	80.1	1	121-364
129	TPR	TPR Domain	4.8e-27	103.3	7	355-388:473-

Table 4

SEQ ID	Pfam Model	Description	E-value	Score	No: of Pfam Domains	Position of the Domain
						506:507- 540:654- 687:688- 721:722- 755:756-789
130 /	HMG14_17	HMG14 and HMG17	1.9e-15	64.7	1	2-73
131	bZIP	bZIP transcription	8.3e-19	71.7	 î	288-352
132	rrm	RNA recognition motif.	1.9e-31	117.9	3	432-502:546- 616:858-929
133	AMP-binding	AMP-binding enzyme	7.1e-117	401.7	1	142-580
138	tubulin	Tubulin/FtsZ family	2.1e-151	516.4	† 1	1-223
141	laminin_EGF	Laminin EGF-like (Domains III and V)	7.6e-12	52.8	4	252-297:300- 348:1342- 1391:1469- 1530
141	Kelch	Kelch motif	1.6e-05	31.8	4	654-702:760- 811:873- 918:929-990
141	integrin_B	Integrins, beta chain	0.0061	9.4	3	44-59:100- 117:1019- 1028
141	EGF	EGF-like domain	0.092	19.3	8	167-203:207- 235:297- 331:496- 533:538- 569:1271- 1308:1312- 1338:1478- 1508
142	RUN	RUN domain	8e-44	159.0	1	31-163
142	FYVE	FYVE zinc finger	2.3e-29	109.1	1	529-593
143	zf-C2H2	Zinc finger, C2H2 type	1.7e-33	124.7	5	442-464:505- 527:533- 555:561- 583:589-611
143	BTB	BTB/POZ domain	1.6e-22	88.2	1	30-143
144	mito_carr	Mitochondrial carrier protein	3.6e-61	216.6	3	10-158:160- 250:254-354
146	DAGKe	Diacylglycerol kinase catalytic domain	0.00015	26.0	1	157-303
147	Exonuclease	Exonuclease	1.6e-41	151.4	1	228-384
147	rrm	RNA recognition motif.	9.5e-08	39.2	2	507-574:602- 674
151	WH2	WH2 motif	6.5e-20	79.6	3	1194- 1214:1234- 1254:1322- 1342
154	DHDPS	Dihydrodipicolinate synthetase family	9.1e-21	82.4	1	3-270
156	PseudoU_synth_1	tRNA pseudouridine synthase	1e-30	115.4	1	111-322
157	pkinase	Protein kinase domain	2.3e-59	210.6	1	216-512
158	ubiquitin	Ubiquitin family	2.4e-05	24.6	1	3-79

Table 4

SEQ	Pfam Model	Description	E-value	Score	No: of	Position of
ID	}				Pfam	the Domain
160	IF-2B	Trisiation fortage 2 and trisia	1.7-00	12402	Domains	155 455
100	H-2B	Initiation factor 2 subunit family	1.7e-98	340.7	1	157-475
161	Beach	Beige/BEACH domain	1.1e-224	759.8	1	1470-1747
161	WD40	WD domain, G-beta repeat	2.9e-08	40.9	5	1848-
			į	1	ţ	1882:1888-
					İ	1928:1947-
			}		1	1983:2030-
			1	1	1	2064:2071-
164	DnaJ	DnaJ domain	1.9e-16	68.1	1	2107
165	Anti_proliferat	BTG1 family	7.4e-85	295.3	1	125-189 11-164
166	sugar_tr	Sugar (and other) transporter	1.2e-78	274.7	1	34-548
167	sugar_tr	Sugar (and other) transporter	7e-52	185.8	1	34-480
168	zf-C2H2	Zinc finger, C2H2 type	1.7e-93	324.0	13	222-244:250-
		Zano iniger, Chill typo	1.,055	324.0	-3	272:278-
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l		1	1			328:334-
1		1		1		356:362-
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}		{	1	1	ł	412:418-
1				1	1	440:446-
		1		1	1	468:474-
ł	l'	Ì	Ì	l	1	496:502-
<u> </u>	,		1			524:530-
168	KRAB	KRAB box	1.8e-35	131.2	1	552:558-580 57-119
169	GBP	Guanylate-binding protein,	1e-191	636.2	11	1-275
		N-terminal domain	10-191		1	19275
169	GBP_C	Guanylate-binding protein, C- terminal domain	6.6e-162	551.3	1	277-573
170	cyclin	Cyclin, N-terminal domain	0.0022	9.3	1	49.100
171	TPR	TPR Domain	9.7e-43	155.4	6	48-192 133-166:167-
1 ***	1118	ITR Domain	9.76-43	133.4	10	200:201-
			l	1	1	234:282-
}]		1	315:316-
		1		1		349:350-383
173	RhoGEF	RhoGEF domain	3.3e-40	147.0	1	166-345
173	PH	PH domain	6.5e-14	54.5	1	378-483
173	SH3	SH3 domain	1.1e-10	48.9	1	72-126
174	zf-C3HC4	Zinc finger, C3HC4 type (RING finger)	0.00011	19.4	1	18-55
174	GBP_C	Guanylate-binding protein, C- terminal domain	0.016	12.1	1	86-114
175	Peptidase_M22	Glycoprotease family	2.3e-73	257.2	1	1-324
177	TBC	TBC domain	4.7e-08	10.1	1	57-268
178	transmembrane4	Tetraspanin family	1.6e-78	259.2	1	16-261
179	CH	Calponin homology (CH) domain	1.2e-25	98.6	1	24-133
179	calponin	Calponin family repeat	1.7e-14	51.8	1	174-199
182	AP_endonucleas1	AP endonuclease family 1	2.6e-17	59.4	2	1-36:50-135
184	Bacterial_PQQ	PQQ enzyme repeat	9.3e-05	29.2	2	52-89:534-
L	1		{	1	1	571

Table 4

SEQ ID	Pfam Model	Description	E-value	Score	No: of Pfam Domains	Position of the Domain
185	DEAD	DEAD/DEAH box helicase	1.6e-60	194.3	1	216-420
185	helicase_C	Helicase conserved C- terminal domain	5.9e-25	96.3	1	454-540
186	zf-C2H2	Zinc finger, C2H2 type	3.2e-24	93.9	6	106-128:134- 156:162- 184:195- 218:477- 499:505-529
187	sugar_tr	Sugar (and other) transporter	0.0014	-90.1	1	272-672
188	tRNA_int_endo	tRNA intron endonuclease, catalytic C-t	0.0025	-7.7	1	73-159
189	WSC	WSC domain	1e-35	132.1	1	175-254
189	Sulfotransfer	Sulfotransferase protein	4e-34	126.8	1	356-586
191	pkinase	Protein kinase domain	5.1e-75	262.6	1	148-421
191	PDZ	PDZ domain (Also known as DHR or GLGF)	1.3e-05	32.1	1	740-827
193	globin	Globin	1.9e-26	96.6	1	3-78
195	WD40	WD domain, G-beta repeat	6.7e-14	59.6	4	64-108:116- 153:158- 194:288-323
197	BRO1	BRO1-like domain	0.0042	-29.4	1	9-161
198	F_actin_cap_B	F-actin capping protein, beta subunit	1.7e-224	759.2	1	1-269
199	ank	Ank repeat	1e-66	235.0	8	40-73:82- 114:115- 147:148- 180:181- 212:213- 246:481- 526:527-559
203	PDZ	PDZ domain (Also known as DHR or GLGF)	4.2e-07	37.0	1	211-293
204	SAM	SAM domain (Sterile alpha motif)	1.2e-11	52.1	1	5-70
205	SAM	SAM domain (Sterile alpha motif)	1.2e-11	52.1	1	5-70
206	zf-UBR1	Putative zinc finger in N- recognin	4.7e-25	96.7	1	978-1046
207	ABC_tran	ABC transporter	2.4e-112	386.6	2	467- 647:1536- 1717
209	zf-C2H2	Zinc finger, C2H2 type	0.00035	27.3	1	200-225
210	UCH-2	Ubiquitin carboxyl-terminal hydrolase family	1.5e-19	78.4	1	385-454
211	IMP4	Domain of unknown function	2.2e-33	124.3	1	144-297
213	zf-C2H2	Zinc finger, C2H2 type	2.9e-08	40.9	3	12-37:173- 198:208-230
214	LysM	LysM domain	2.1e-11	51.3	1	73-116
215	ank	Ank repeat	1.1e-05	32.3	2	834-867:879- 912
215	TIG	IPT/TIG domain	0.009	22.6	1	642-723
217	pyr_redox	Pyridine nucleotide-	1.7e-71	251.0	1	196-470

Table 4

SEQ ID	Pfam Model	Description	E-value	Score	No: of Pfam Domains	Position of the Domain
		disulphide oxidoreducta				
217	Rieske	Rieske [2Fe-2S] domain	6.2e-20	79.6	1	68-168
218	PDZ	PDZ domain (Also known as DHR or GLGF)	8.5e-19	75.9	1	642-728
219	pkinase	Protein kinase domain	8.1e-67	235.4	1	26-204
220	dsrm	Double-stranded RNA binding motif	0.095	7.5	1	100-172
221	PHD	PHD-finger	5.4e-05	29.6	1	147-203
222	L27	L27 domain	6.5e-16	66.3	1	13-68
222	SAM	SAM domain (Sterile alpha motif)	7.2e-10	46.2	2	1051- 1117:1166- 1230
223	TRM	N2,N2-dimethylguanosine tRNA methyltransfera	7.3e-22	86.1	1	227-693
224	LIM	LIM domain	5.3e-06	33.4	2	124-180:183- 243
225	ig	Immunoglobulin domain	1.1e-07	29.8	1	55-144
227	F-box	F-box domain	1.3e-05	32.1	1	11-59
229	Glucosamine_iso	Glucosamine-6-phosphate isomerases/6-	2.7e-158	539.3	1	15-250
231	PTN_MK	PTN/MK heparin-binding protein family	3.6e-44	160.2	1	51-148
236	ion_trans	Ion transport protein	1.6e-22	88.3	1	174-393
238	GNS1_SUR4	GNS1/SUR4 family	5.2e-46	166.3	1	10-265
240	ubiquitin	Ubiquitin family	2.7e-05	24.4	1	10-89
241	PIP5K	Phosphatidylinositol-4- phosphate 5-Kinase	1.5e-155	530.2	1	124-420
242	cadherin	Cadherin domain	0	1298.9	19	1-75:89- 180:194- 290:355- 434:448- 549:563- 652:671- 774:788- 881:896- 988:1002- 1092:1106- 1192:1206- 1295:1309- 1379:1393- 1489:1503- 1594:1608- 1699:1713- 1808:1814- 1910:1922- 2016
244	fn3	Fibronectin type III domain	1.2e-31	118.6	4	58-140:152- 238:249- 333:345-426
245	UQ_con	Ubiquitin-conjugating enzyme	1.4e-16	68.5	1	93-250
246	LRR	Leucine Rich Repeat	1.7e-14	61.6	6	51-75:76-

Table 4

SEQ ID	Pfam Model	Description	E-value	Score	No: of Pfam Domains	Position of the Domain
						99:155- 178:181- 203:204- 226:227-251
247	lipocalin	Lipocalin / cytosolic fatty- acid binding pr	1.2e-28	102.8	1	164-294
248	Ribosomal_S2	Ribosomal protein S2	2.9e-11	43.7	1	33-80
249	tubulin	Tubulin/FtsZ family	8.5e-163	554.2	1	1-277
250_	tubulin	Tubulin/FtsZ family	2.4e-212	718.8	1	1-351
251	ATP-synt_ab	ATP synthase alpha/beta family, nucleot	1.2e-75	264.8	1	138-346
251	ATP-synt_ab_C	ATP synthase alpha/beta chain, C termin	2.7e-38	140.6	1	348-456
251	ATP-synt_ab_N	ATP synthase alpha/beta family, beta-ba	5.4e-19	76.5	1	67-135
252	ATP-synt_ab	ATP synthase alpha/beta family, nucleot	1.3e-70	248.0	1	138-344
252	ATP-synt_ab_N	ATP synthase alpha/beta family, beta-ba	5.4e-19	76.5	1	67-135
253	zf-C3HC4	Zinc finger, C3HC4 type (RING finger)	5e-12	43.2	1	39-79
254	G-patch	G-patch domain	1.3e-08	42.1	1	410-456
255	CH	Calponin homology (CH) domain	1.6e-11	51.7	1	24-134
256	RF-1	Peptidyl-tRNA hydrolase domain	5.9e-66	232.5	1	225-338
257	RF-1	Peptidyl-tRNA hydrolase domain	5.9e-66	232.5	1	189-302
258	OTU	OTU-like cysteine protease	4.4e-18	73.5	1	189-304
259	thiored	Thioredoxin	2e-09	35.7	2	119-165:662- 695
260	thyroglobulin_1	Thyroglobulin type-1 repeat	3.1e-34	127.2	2	95-158:227- 292
260	kazal	Kazal-type serine protease inhibitor	9.36-07	35.9	1	43-87
262	DnaJ	DnaJ domain	4.1e-15	63.6	1	277-338
263	WD40	WD domain, G-beta repeat	4e-21	83.6	5	3-42:49- 86:97- 133:142- 178:184-220
265	DUF6	Integral membrane protein DUF6	0.083	9.1	2	81-316:338- 470
266	Ribosomal_L31e	Ribosomal protein L31e	1.7e-61	217.7	1	15-109
268	F5_F8_type_C	F5/8 type C domain	2.4e-65	230.5	1	42-196
268	Zn_carbOpept	Zinc carboxypeptidase	3.5e-50	180.1	2	224-341:400- 600
270	BTB	BTB/POZ domain	7.7e-18	72.7	1	8-119
270	zf-C2H2	Zinc finger, C2H2 type	4.2e-13	57.0	4	254-276:363- 385:390- 412:448-468
271	Glycos_transf_1	Glycosyl transferases group 1	0.027	12.8	1	291-385
272	HEAT	HEAT repeat	2.2e-07	38.0	3	237-275:276-

Table 4

SEQ	Pfam Model	Description	E-value	Score	No: of	Position of
ID					Pfam Domains	the Domain
						315:674-712
273	HEAT	HEAT repeat	2.2e-07	38.0	3	237-275:276-
			l	<u> </u>		315:640-678
275	SPRY	SPRY domain	2.6e-34	127.4	1	390-515
275	zf-C3HC4	Zinc finger, C3HC4 type (RING finger)	1e-16	58.5	1	29-69
277	BTB	BTB/POZ domain	6e-27	103.0	1	36-149
277	Kelch	Kelch motif	9.7e-21	82.3	4	331-390:392-
))			441:443-
				 		493:540-586
278	zf-C2H2	Zinc finger, C2H2 type	4.1e-116	399.2	14	193-215:221-
			{	}		243:249-
•				1	· ·	271:277-
]	Ì		1]		299:305- 327:333-
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		· ·		ł		439:445-
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•					ĺ	495:501-
		{		ł	}	523:529-
}			1	1	1	551:557-579
229	SCAN	SCAN domain	2.4e-52	187.3	1	36-132
229	zf-C2H2	Zinc finger, C2H2 type	2.4e-51	184.0	7	348-370:375-
(į			Ì		397:403-
1			ì	ì	1	425:431-
ļ]		1]	1	453:459-
]		į	1]]	480:486-
		CONTRACT.	1	70.6		508:514-537
231	Zip	ZIP Zinc transporter	6.6e-20	79.6	1	1-146
282	NTP_transf_2	Nucleotidyltransferase domain	8.5e-13	55.9	1	67-174
286	zf-C2H2	Zinc finger, C2H2 type	2.8e-93	323.3	12	118-140:146-
1	}			1	1	168:174-
}		İ		1		196:202-
				1.		224:230-
		1	1		{	252:258-
		ł	1	1	1	280:286-
ļ		}			1	308:314-
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				į .		364:370- 392:398-
l	1	4		1	1 .	420:426-448
286	KRAB	KRAB box	3.6e-38	140.2	1	8-70
287	zf-C2H2	Zinc finger, C2H2 type	5.3e-124	425.4	17	183-205:211-
1 -0'		- ingo, care type	3.30-127		1 ~	233:239-
1	1					261:267-
ł			l		1	289:295-
)		}]]	,	317:323-
1						345:351-
			1		Í	373:379-

Table 4

SEQ ID	Pfam Model	Description	E-value	Score	No: of Pfam Domains	Position of the Domain
						401:407- 429:435-
	1		j	1	}	457:463-
	· ·		l	1	} .	485:491-
	ļ		ł	Ì		513:519-
	1					541:547-
	· · ·		ì	ł		569:575-
			Į.	ļ	ļ	597:603-
		· · · · · · · · · · · · · · · · · · ·	1	<u> </u>	<u> </u>	625:631-653
289	DiHfolate_red	Dihydrofolate reductase	7.4e-77	268.8	1	4-185
291	PDZ	PDZ domain (Also known as DHR or GLGF)	7.4e-17	69.4	1	5-84
293	PH	PH domain	1.4e-08	35.5	1	44-147
294	adh_short	short chain dehydrogenase	3.9e-29	110.2	1	36-284
297	PKD	PKD domain	9.9e-09	42.4	2	663-753:756- 839
297	BNR	BNR repeat	3.2e-06	34.1	5	115-126:156- 167:351- 362:428- 439:470-481
300	HMG_box	HMG (high mobility group) box	5.4e-05	20.0	1	245-304
301	ig	Immunoglobulin domain	0.05	11.6	1	629-688
302	zf-C3HC4	Zinc finger, C3HC4 type (RING finger)	5e-12	43.2	1	39-79
303	START	START domain	0.015	4.1	1	1790-1994
304	integrase	Integrase DNA binding domain	7.2e-06	32.9	1	51-96
305	myosin_head	Myosin head (motor domain)	7.6e-279	939.7	2 .	11-668:689- 733
306	zf-C2H2	Zinc finger, C2H2 type	8.5e-54	192.1	7	66-88:94- 116:122- 144:150- 172:178- 200:280- 303:317-339
307	ig	Immunoglobulin domain	0.00023	19.1	2	35-104:136- 194
309	ras	Ras family	0.00079	-93.3	1	38-176
310	ig	Immunoglobulin domain	2.1e-06	25.7	1	37-112
311	EFIBD	EF-1 guanine nucleotide exchange domain	4.7e-56	199.6	1	139-225
312	BTB	BTB/POZ domain	8.4e-25	95.8	1	51-164
313	zf-C2H2	Zinc finger, C2H2 type	7.76-59	208.9	9	118-140:197- 219:281-
						303:309- 331:337- 359:365- 387:393- 415:421-
	<u></u>				<u></u>	443:449-471
313	KRAB	KRAB box	1.4e-17	71.8	1	41-99

Table 4

SEQ ID	Pfam Model	Description	E-value	Score	No: of Pfam Domains	Position of the Domain
314	Hydrolase	haloacid dehalogenase-like hydrolase	0.045	8.2	1	213-671
315	cNMP_binding	Cyclic nucleotide-binding domain	4e-26	100.2	1	387-475
315	ion_trans	Ion transport protein	3.8e-19	77.0	1	69-290
316	Peptidase_S26	Signal peptidase I	2.8e-16	56.3	2	38-98:117- 139
317	zf-C2H2	Zinc finger, C2H2 type	4.3e-56	199.8	9	156-178:184- 206:212- 234:240- 262:268- 290:296- 318:324- 346:352- 374:378-400
317	KRAB	KRAB box	6.7e-16	66.3	1	11-73
319	UPF0073	Uncharacterised protein family	1.8e-09	27.9	1	33-276
320	EGF	EGF-like domain	4.7e-08	40.2	1	26-59
321	lectin_c	Lectin C-type domain	8.6e-15	62.6	1	268-374
325	MAM	MAM domain	1.3e-52	188.2	1	338-503
325	ig	Immunoglobulin domain	1.9e-15	54.8	3	41-101:138- 202:346-420
327	MAM	MAM domain	5.3e-180	611.4	4	26-169:170- 329:342- 498:509-666
328	Sema	Sema domain	1.5e-211	716.2	1	56-491
329	zf-C2H2	Zinc finger, C2H2 type PAP2 superfamily	1.5e-84	294.3	13	170-192:198- 220:226- 248:254- 276:282- 304:310- 332:338- 360:366- 388:394- 416:422- 444:450- 472:478- 500:506-528
332	LRR	Leucine Rich Repeat	3.4e-36	133.7	11	58-81:82-
332		Leating Rich Repeat	3,40-30	133./		105:106- 129:130- 153:154- 177:178- 201:202- 225:250- 273:274- 297:298- 321:322-345
332	ig	Immunoglobulin domain	2.5e-08	31.9	1	425-485
332	LRRNT	Leucine rich repeat N-	2.5e-05	31.1	1	27-56

Table 4

SEQ ID	Pfam Model	Description	E-value	Score	No: of Pfam Domains	Position of the Domain
		terminal domain				
332	LRRCT	Leucine rich repeat C- terminal domain	0.0029	24.3	1	355-408
333	AdoHcyase	S-adenosyl-L-homocysteine hydrolase	1.5e-280	945.4	1	214-640
334	TBC	TBC domain	9.4e-38	138.9	1	89-302
341	WD40	WD domain, G-beta repeat	0.00094	25.9	2	2-32:109-146
342	ABC1	ABC1 family	0.051	-29.9	1	3-50
344	globin	Globin	3e-45	162.2	1	1-141
345	globin	Globin	7.5e-39	139.9	2	1-31:68-179
347	F-box	F-box domain	1.5e-07	38.5	1	24-72
348	HLH	Helix-loop-helix DNA- binding domain	2e-08	41.4	i	83-137
349	KRAB	KRAB box	2.7e-39	144.0	1	4-66
350	UCH-2	Ubiquitin carboxyl-terminal hydrolase family	1.7e-19	78.2	1	645-705
350	UCH-1	Ubiquitin carboxyl-terminal hydrolases famil	9.1e-15	62.5	1	363-394
350	zf-UBP	Zn-finger in ubiquitin- hydrolases and other	0.00069	18.9	1	236-306
351	NUDIX	MutT-like domain	8.2e-12	52.7	1	50-200
352	IBR	IBR domain	1.6e-12	55.0	1	101-166
353	IBR	IBR domain	1.6e-12	55.0	1	66-131
354	SCP	SCP-like extracellular protein	1.4e-34	128.3	1	56-208
356	mito_carr	Mitochondrial carrier protein	9.7e-78	271.7	3	10-125:127- 220:232-321
358	UCH-1	Ubiquitin carboxyl-terminal hydrolases famil	5.1e-15	63.3	1	323-354
358	zf-UBP	Zn-finger in ubiquitin- hydrolases and other	0.00049	19.4	1	195-264
360	Phage_lysozyme	Phage lysozyme	0.0014	23.4	1	94-184
362	Ribosomal_S2	Ribosomal protein S2	3.3e-08	32.9	1	20-62
364	zf-C3HC4	Zinc finger, C3HC4 type (RING finger)	5.3e-09	33.4	1	291-329
365	zf-C3HC4	Zinc finger, C3HC4 type (RING finger)	0.0096	13.1	1	109-148
367	TPR	TPR Domain	0.043	20.4	1	1-28
370	zf-C2H2	Zinc finger, C2H2 type	5.3e-109	375.5	14	127-149:155- 177:183- 205:211- 233:239-
						261:267- 289:295- 317:323-
	_					345:351- 373:379- 401:407- 429:435- 457:463- 485:491-513
370	SCAN	SCAN domain	4.2e-38	140.0	1	27-122
371	arf	ADP-ribosylation factor	4.9e-39	143.1	1	6-184

Table 4

SEQ ID	Pfam Model	Description .	E-value	Score	No: of Pfam Domains	Position of the Domain
		family		1		
371	ras	Ras family	7.2e-06	-70.1	1	22-186
372	BNR	BNR repeat	0.031	20.9	3	171-182:244- 255:295-306
373	zf-C2H2	Zinc finger, C2H2 type	8.3e-25	95.8	5	142-162:171- 198:204- 228:234- 258:264-288
376	rm	RNA recognition motif.	0.00019	28.2	11	112-163
377	rrm	RNA recognition motif.	2.2e-19	77.9	1	112-183
380	vwc	von Willebrand factor type C domain	1.6e-31	118.2	3	22-76:79- 134:137-192
381	Ribosomal_L35Ae	Ribosomal protein L35Ae	0.00013	7.0	1	1-79
385	ras	Ras family	3.9e-63	223.2	1	35-229
385	arf	ADP-ribosylation factor family	1.7e-05	-46.9	1	18-202
388	F-box	F-box domain	1.5e-05	31.9	2	23-70:99-146
390	SPRY	SPRY domain	6.2e-10	46.4	1	101-239
391	tRNA_Me_trans	tRNA methyl transferase	1.9e-19	50.9	1	5-185
392	zf-C2H2	Zinc finger, C2H2 type	4e-17	70.3	3	175-197:203- 225:231-253
393	SCAN	SCAN domain	3.1e-39	143.8	1	389-484
393	SPRY	SPRY domain	1.8e-19	78.1	1	148-273
393	zf-C2H2	Zinc finger, C2H2 type	4e-09	43.7	2	759-781:787- 809
393	zf-C3HC4	Zinc finger, C3HC4 type (RING finger)	0.0032	14.7	1	11-52
394	Kelch	Kelch motif	4e-53	189.9	5	329-375:377- 431:433- 479:481- 525:527-572
394	BTB	BTB/POZ domain	6.1e-26	99.6	1	30-144
395	C2	C2 domain	2.2e-80	280.4	2	159-251:296- 384
396	ank	Ank repeat	5.6e-33	123.0	4	47-79:80- 112:140- 174:175-207
396	PH	PH domain	8.9e-05	22.0	1	236-334
397	ank	Ank repeat	1.7e-26	101.4	4	17-49:50- 82:83- 115:116-148
398	Nucleoplasmin	Nucleoplasmin	3.6e-29	110.4	1	13-209
400	DAGKa	Diacylglycerol kinase accessory domain	1.9e-124	426.8	1	598-778
400	DAGKc	Diacylglycerol kinase catalytic domain	7.1e-67	235.6	1	454-578
400	DAG_PE-bind	Phorbol esters/diacylglycerol binding dom	2.9e-23	90.7	2	261-310:326- 374
400	efhand	EF hand	2.4e-12	54.4	2	169-197:214- 242
403	PDZ	PDZ domain (Also known as	7.7e-46	165.7	3	86-166:210-

Table 4

SEQ ID	Pfam Model	Description	E-value	Score	No: of Pfam Domains	Position of the Domain
		DHR or GLGF)				291:821-907
404	zf-C2H2	Zinc finger, C2H2 type	2.6e-48	173.9	7	172-194:200- 222:228- 250:256- 278:284- 306:312-
405	K_tetra	K+ channel tetramerisation domain	2.6e-23	90.9	1	331:340-362 51-146
406	SNF	Sodium:neurotransmitter symporter family	0	1268.7	1	60-657
407	ig	Immunoglobulin domain	1.1e-06	26.5	1	53-120
408	DnaJ	DnaJ domain	2.3e-27	104.3	1	4-68
408	DnaJ_C	DnaJ C terminal region	3.1e-08	38.1	1	192-314
409	mito_carr	Mitochondrial carrier protein	1.4e-57	204.7	3	5-100:102- 201:205-302
410	zf-C2H2	Zinc finger, C2H2 type	5.2e-97	335.7	12	141-163:169- 191:197- 219:225- 247:253- 275:281- 303:309- 331:337- 359:365-
						387:393- 415:421- 443:449-473
411	S_100	S-100/ICaBP type calcium binding domain	9.7e-13	55.8	. 1	5-48
411	efhand	EF hand	0.0012	25.6	1	54-82
413	fn3	Fibronectin type III domain	8.6e-14	59.3	2	22-107:119- 196
413	PHD	PHD-finger	9.6e-05	27.2	1	285-341
414	zf-C2H2	Zinc finger, C2H2 type	2.3e-27	104.4	6	42-64:117- 139:145- 167:173- 196:534- 556:573-595
415	SPRY	SPRY domain	3.9e-18	73.7	1	347-467
415	zf-C3HC4	Zinc finger, C3HC4 type (RING finger)	4.4e-14	49.9	1	16-56
415	zf-B_box	B-box zinc finger	9e-07	35.9	1	92-133
416	pkinase	Protein kinase domain	1.2e-54	195.0	1	97-317
417	trypsin	Trypsin	4.6e-38	122.5	1	41-234
418	Glypican Keratin_B2	Glypican Keratin, high sulfur B2 protein	5.7e-131 0.0013	-23.4	1	3-244 37-159
420	Dynein_heavy	Dynein heavy chain	0	1432.3	† <u>1</u>	309-1019
421	zf-C2H2	Zinc finger, C2H2 type	0.00039	27.2	3	75-99:203- 227:266-290
422	ig	Immunoglobulin domain	0.00074	17.5	1	34-107
423	fn3	Fibronectin type III domain	6e-08	39.8	1	443-531

Table 4

SEQ ID	Pfam Model	Description	E-value	Score	No: of Pfam Domains	Position of the Domain
424	Keratin_B2	Keratin, high sulfur B2 protein	0.0023	-27.1	2	5-150:152- 251
425	pkinase	Protein kinase domain	2.3e-55	197.3	1	69-390
426	ig	Immunoglobulin domain	4.1e-09	34.4	1	35-112
427	Galactosyl_T	Galactosyltransferase	2.6e-35	130.8	1	158-349
428	proteasome	Proteasome A-type and B- type	5.5e-28	106.4	1	96-238
429	7tm_1	7 transmembrane receptor (rhodopsin family)	3.4e-38	123.5	1	41-290
430	BTB	BTB/POZ domain	8.1e-23	89.2	1	58-173
430	zf-C2H2	Zinc finger, C2H2 type	4.3e-07	37.0	2	472-494:500- 523
433	p450	Cytochrome P450	6.4e-175	594.5	1	33-493
434	sugar_tr	Sugar (and other) transporter	2.6e-64	227.1	1	10-512
435	zf-C2H2	Zinc finger, C2H2 type	1.8e-52	187.8	9	287-309:315- 337:546- 568:574- 596:606- 628:844- 866:872- 894:980- 1002:1008- 1030
436	7tm_1	7 transmembrane receptor (rhodopsin family)	2.2e-40	130.4	2	82-221:229- 284
437	FGF	Fibroblast growth factor	4.6e-14	51.6	1	48-129
438	Osteopontin	Osteopontin	3.7e-181	615.2	1	1-294

							Pre-street about			Personal Control
PDB annotation	TRANSCRIPTION REGULATION PROTO- ONCOGENE, NUCLEAR BODIES (PODS), LEUKEMIA, 2 TRANSCRIPTION REGULATION	TRANSFERASE HRS, HRS, VHS, FYVE, ZINC FINGER, SIPPRHFI IX	METAL BINDING PROTEIN RING FINGER PROTEIN MATI; RING FINGER (C3HC4)	COMPLEX (ISOMERASE/DIPEPTIDE) PIN1; PEPTIDYL-PROLYL CIS-TRANS ISOMERASE, ROTAMASE, 2 COMPLEX (ISOMERASE/DIPEPTIDE) CONECT		COMPLEX (APOPTOSIS/PEPTIDE) APOPTOSIS, ALTERNATIVE SPLICING, COMPLEX (APOPTOSIS/PEPTIDE)	APOPTOSIS HELICAL (I	APOPTOSIS APOPTOSIS REGULATOR BCL-X; APOPTOSIS, PROGRAMMED CELL DRATH, BCL-2	5,00	STRUCTURAL PROTEIN II
Compound	TRANSCRIPTION FACTOR PML; CHAIN: NULL;	HEPATOCYTE GROWTH FACTOR-REGULATED TYROSINE CHAIN: A:	CDK-ACTIVATING KINASE ASSEMBLY FACTOR MAT1; CHAIN: A;	PEPTIDYL-PROLYL CIS- TRANS ISOMERASE; CHAIN: A; ALA-PRO DIPEPTIDE; CHAIN: B;		BCL-XL; CHAIN: A; BAK PEPTIDE; CHAIN: B;	APOPTOSIS REGULATOR BAX, MEMBRANE ISOFORM ALPHA; CHAIN: A:	BCL-XL; CHAIN: NULL;		ALPHA SPECTRIN; CHAIN:
SEQ FOLD score										81.85
PMF score	0.01	0.76	0.62	0.23		0.01	0.10	0.03		
Verify score	-0.58	-0.23	-0.44	-0.45		-0.07	-0.28	-0.03		
Psi Blast	5.2e-09	19000	1.3e-06	7.8e-06		5.1e-38	1.7e-36	3.46-40		1e-21
END AA	09	72	09	98		362	362	362		297
START AA	01	2	10	8		223	208	223		94
CHAIN		A	¥	∢		4	Ą			A
PDB ID	Ibor	ldvp	1825	1pin		Ibxl	1f16	lmaz		lcun
SEQ ID NO:		1	-	vo	·	11	11	11		12

\Box	1		Т	· T											Th.	7	i ik	Harry.	le H	- Rendi	,	- IE	E 4			
PDB annotation	TWO REPEATS OF SPECTRIN, ALPHA HELICAL LINKER REGION, 2 2 TANDEM 3-HELIX COLLED- COLLS, STRUCTURAL	CONTRACTLE PROTEIN TRIPLE-HELIX COILED COIL, CONTRACTILE PROTEIN		RIBOSOME 50S RIBOSOMAL	50S RIBOSOMAL PROTEIN	L3P, HIMAL3, HL1; 50S	KIBOSOMAL PROTEIN LAE, HMAI 4 HI 6: 50S	RIBOSOMAL PROTEIN LSP.	HMAL5, HL13; 30S	RIBOSOMAL PROTEIN HS6;	113P HMAI 13: 508	RIBOSOMAL PROTEIN L14P,	HMAL14, HL27; 50S	RIBOSOMAL PROTEIN LISP	HMALIS, HL9; 50S	HMAL18. HL12: 50S	RIBOSOMAL PROTEIN L18E	HI 29, L19; 50S RIBOSOMAL	PROTEIN L19E, HMAL19,	HL24; 50S RIBOSOMAL	PROTEIN L21E, HL31; 50S	RIBOSOMAL PROTRIN L22P.	HMAL22, HL23; 50S RIBOSOMAI PROTERNI 23P	HMAL23, HL25, L21: 50S	4P,	HMAL24, HL16, HL15; 50S
Compound	A, B, C;	HUMAN SKELETAL MUSCLE ALPHA-ACTININ 2; CHAIN: A;		23S RRNA; CHAIN: 0; 5S	RIBOSOMAL PROTEIN L2;	CHAIN: A; RIBOSOMAL	RIBOSOMAL PROTEIN 14:	CHAIN: C; RIBOSOMAL	PROTEIN LS; CHAIN: D;	RIBOSOMAL PROTEIN	RIBOSOMAI, PROTEIN	L10E; CHAIN: F;	RIBOSOMAL PROTEIN	L13; CHAIN: G;	KIBOSOMAL PROTEIN	RIBOSOMAL PROTEIN	LISE; CHAIN: I;	RIBOSOMAL PROTEIN	L15; CHAIN: J;	RIBOSOMAL PROTEIN	LIS; CHAIN: K;	KIBOSOMAL PROTEIN	LISE; CHAIN: L; RIBOSOMAI, PROTRIN	L19; CHAIN: M;	RIBOSOMAL PROTEIN	L21E; CHAIN: N;
SEQ FOLD score		72.83									• •															
PMF	·_			0.00																					-	
Verify score				-0.90										•												
Psi Blast		3.4e-11		2.6e-26																						
END AA		297		142																			•		:	
START AA		77		75	 -		•	-	-						•				•							
CHAIN		Ą		Þ				-											-			•				
PDB ID		Iquu		1旗																	-					
SEQ ID NO:		12		14																						

Table :

						7)														17-17	· ·	ĘC.		12		11 11		-	11 11	न्या			
PDB annotation	RIBOSOMAL PROTEIN L24E, HL21/HL22; 50S RIBOSOMAL	PROTEIN L29P, HMAL29, HT 33- 50S PIROSOMAT	PROTEIN L30P, HIMAL30,	HL20, HL16; 50S	RIBOSOMAL PROTEIN L31E,	L34, HL30; 50S RIBOSOMAL	PROTEIN LIZE, HLS; 30S	KIBOSOMAL FROI EIN L3/E,	PROTEINS L39E, HL39E,	HL46E; 50S RIBOSOMAL	PROTEIN LAME, LA, HLA; 50S	RIBOSOMAL PROTEIN LOP,	HMAL6, HL10 RIBOSOME	ASSEMBLY, RNA-RNA,	PROTEIN-KINA, FROIEBY-	INGIER					li-e-	RIBOSOME 50S RIBOSOMAL	PROTEIN L2P, HMAL2, HL4;	50S RIBOSOMAL PROTEIN	L3P, HMAL3, HL1; 50S	RIBOSOMAL PROTEIN LAE,	HMAL4, HL6; 508	RIBOSOMAL PROTEIN LSP, C.	DIBOSONAL DEPOTERNIES	SOS RIBOSOMAI, PROTEIN	L13P, HMAL13; 50S	RIBOSOMAL PROTEIN L14P	HMAL14, FILL 1; 505
Compound	RIBOSOMAL PROTEIN L22; CHAIN: 0;	RIBOSOMAL PROTEIN	RIBOSOMAL PROTEIN	1.24; CHAIN: Q;	RIBOSOMAL PROTEIN	L24E; CHAIN: R;	KIBOSOMAL PROTEIN	RIBOSOMAL PROTEIN	L30: CHAIN: T:	RIBOSOMAL PROTEIN	L31E; CHAIN: U;	RIBOSOMAL PROTEIN	L32E; CHAIN: V;	RIBOSOMAL PROTEIN	L3/AE; CHAMN: W;	1.37E: CHAIN: X:	RIBOSOMAL PROTEIN	L39E; CHAIN: Y;	RIBOSOMAL PROTEIN	L44E; CHAIN: Z;	RIBOSOMAL PROTEIN L6;	23S RRNA; CHAIN: 0; 5S	RRNA; CHAIN: 9;	RIBOSOMAL PROTEIN L2;	CHAIN: A; RIBOSOMAL	PROTEIN L3; CHAIN: B;	RIBOSOMAL PROTEIN L4;	CHAIN: C; KIBOSOMAL	PIBOCOMAI DECIPEN	TAR: CHAIN: F.	RIBOSOMAL PROTEIN	L10E; CHAIN: F;	KIBUSUMAL PROTEIN
SEQ FOLD score																																	
PMF score											`										·	0.12											
Verify				•												•						-0.40											
Psi Blast																					•	5.1e-21										•	
END																						152											
START AA																						78											
CHAIN															_						_	þ										- 	7
PDB CD																						1ffk											
SEQ ID NO:																						4											

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PDB annotation	RIBOSOMAL PROTEIN LISP, HMAL15, HL9; 50S	RIBOSOMAL PROTEIN L18P,	HMAL18, HL12; 50S	RIBOSOMAL PROTEIN LISE,	PROTEIN I 19R HWAI 19	HL24; 50S RIBOSOMAL	PROTEIN L21E, HL31; 50S	RIBOSOMAL PROTEIN L22P,	HMAL22, HL23; 50S	RIBOSOMAL PROTEIN L23P,	HMAL23, HL25, L21; 50S	HMAI 24 HI 16 HI 15: 508	RIBOSOMAL PROTEIN 1.24E.	HL21/HL22; 50S RIBOSOMAL	PROTEIN L29P, HMAL29,	HL33; 50S RIBOSOMAL	PROTEIN L30P, HMAL30,	HL20, HL16; 50S	RIBOSOMAL PROTEIN L31E,	L34, HL30; 50S RIBOSOMAL	PROTEIN L32E, HLS; 50S	TASE SON RIBONOMAT	PROTEINS L39E, HL39E.	HI 46E; 50S RIBOSOMAL	PROTEIN LAGE, LA, HLA; 505	RIBOSOMAL PROTEIN LGP, IJ	HMAL6, HL10 RIBOSOME	ASSEMBLY, RNA-RNA,	PROTEIN-RNA, PROTEIN-	PROTEIN	!		The Change	
Compound	L13; CHAIN: G; RIBOSOMAL PROTEIN	L14; CHAIN: H;	RIBOSOMAL PROTEIN	LISE; CHAIN: 1; DIBOSOMAT BROTTEIN	L15: CHAIN: 1:	RIBOSOMAL PROTEIN	L18; CHAIN: K;	RIBOSOMAL PROTEIN	L18E; CHAIN: L;	RIBOSOMAL PROTEIN	PIBOSOMAI PROTEIN	L21E: CHAIN: N:	RIBOSOMAL PROTEIN	L22; CHAIN: 0;	RIBOSOMAL PROTEIN	L23; CHAIN: P;	RIBOSOMAL PROTEIN	L24; CHAIN: Q;	RIBOSOMAL PROTEIN	L'24E; CHAIN: K;	KIBOSOMAL PROTEIN	RIBOSOMAI, PROTRIN	L30; CHAIN: T;	RIBOSOMAL PROTEIN	L31B; CHAIN: U;	RIBOSOMAL PROTEIN	L32E; CHAIN: V;	RIBOSOMAL PROTEIN	L37AE; CHAIN: W;	KIBOSOMAL PROTEIN	L3/E; CHAIN: X;	L39E; CHAIN: Y:	RIBOSOMAL PROTEIN	L44E; CHAIN: Z;
SEQ FOLD score									·					·															-					
PMF score									•								•									-								
Verify							,	æ																						-				
Psi Blast						,																···	·						•			•		1
END																											_						•	1
START AA																						l					•							
CHAIN ID																		-					-							٠	-			T
PDB ID																																		1
SEQ ID NO:																																		

				τ-	·		gran.	Free adm	स्य	tot leadt andr De Heal	E 445	- TP	hand E.
PDB annotation		CHAPERONE HOP, TPR- DOMAIN, PEPTIDE- COMPLEX, HELICAL REPEAT, HSP90, 2 PROTEIN BINDING	SIGNALING PROTEIN PEROXISMORE RECEPTOR 1, PTS1-BP, PEROXIN-5, PTS1 PROTEIN-PEPTIDE COMPLEX, TETRATRICOPEPTIDE REPEAT, TPR, 2 HELICAL REPEAT		LYASE EPIMERASE, DEHYDRATASE, DEHYDROGENASE, LYASE	ISOMERASE EPIMERASE; UDP-GALACTOSE, EPIMERASE, ISOMERASE		TRANSFERASE MURG; ROSSMANN FOLD	23	AMINOACYL-TRNA SYNTHETASE METRS; AMINOACYL-TRNA SYNTHETASE, ROSSMANN FOLD	AMINOACYL-TRNA KANTHASE		
Compound	RIBOSOMAL PROTBIN L6; CHAIN: 1;	TPRZA-DOMAIN OF HOP; CHAIN: A; HSP90-PEPTIDE MEEVD; CHAIN: B;	PEROXISOMAL TARGETING SIGNAL 1 RECEPTOR; CHAIN: A, B; PTS1-CONTAINING PEPTIDB; CHAIN: C, D;		DTDP-GLUCOSE 4,6- DEHYDRATASE; CHAIN: A. B:	UDP-GALACTOSE4- EPIMERASE; CHAIN: NULL;		UDP-N- ACETYLGLUCOSAMINE- N-ACETYLMURAMYL- CHAIN: A, B;		METHIONYL-TRNA SYNTHETASB; CHAIN: NULL;	GLUTAMYL-TRNA SYNTHETASE; 1GLN 4 CHAIN: NULL 1GLN 5		VIRUS EQUINE HERPES
SEQ FOLD score	·				55.95	72.32				103.94	74.49		
PMF score		-0.15	0.04					0,59			·		0.95
Verify		0.04	-0.24					0.19		-			0.18
Psi Blast		3.4e-11	5.le-17		1e-72	5.1e-63		5.1e-20		1.7e-57	8.5e-40		5.1e-11
END		420	450		361	363		410		563	537		76
START AA	•	322	174		11	11		208		71	73		33
CHAIN		А	V		₹			Ą			-		
PDB CD		lelr	1fch		1bxk	1udb		1f0k		1a8h	1gln		1chc
SEQ ID NO:			16		17	17		25		30	30		32

Table 5

			•		1000	11 11 11 11 11 11 11 11 11 11 11 11 11	
PDB annotation		LIGASE CBL, UBCH7, ZAP- 70, E2, UBIQUITIN, E3, PHOSPHORYLATION, 2 TYROSINE KINASE, UBIQUITINATION, PROTEIN DEGRADATION,	LIGASE CBL, UBCH7, ZAP- 70, E2, UBIQUITIN, E3, PHOSPHORYLATION, 2 TYROSINE KINASE, UBIQUITINATION, PROTEIN DEGRADATION,	METAL BINDING PROTEIN RING FINGER PROTEIN MAT1; RING FINGER (C3HC4)	METAL BINDING PROTEIN RING FINGER PROTEIN MAT1: RING FINGER (C3HC4)	DNA-BINDING PROTEIN V(D)J RECOMBINATION ACTIVATING PROTEIN 1; RAG1, V(D)J RECOMBINATION, ANTIBODY, MAD, RING FINGER, 2 ZINC BINUCLEAR CLUSTER, ZINC FINGER, DNA-BINDING PROTEIN	DNA-BINDING PROTEIN (VD)J RECOMBINATION ACTIVATING PROTEIN 1; PRAG1, V(D)J RECOMBINATION, RECOMB
Compound	VIRUS-1 (C3HC4, OR RING DOMAIN) 1CHC 3 (NMR, 1 STRUCTURE) 1CHC 4	SIGNAL TRANSDUCTION PROTEIN CBL; CHAIN: A; ZAP-70 PEPTIDE; CHAIN: B; UBIQUITIN- CONTUGATING ENZYME B12-18 KDA UBCH7; CHAIN: C.	SIGNAL TRANSDUCTION PROTEIN CBL; CHAIN: A; ZAP-70 PEPTIDE; CHAIN: B; UBIQUITIN- CONTUGATING ENZYME E12-18 KDA UBCH7; CHAIN: C.	CDK-ACTIVATING KINASE ASSEMBLY FACTOR MAT1; CHAIN: A;	CDK-ACTIVATING KINASE ASSEMBLY FACTOR MAT1; CHAIN: A;	RAGI; CHAIN: NUIL;	RAGI; CHAIN: NULL;
SEQ FOLD score	·					·	60.57
PMF score	·	0.33	0.93	0.25	0.51	1.00	
Verify score	,	0.12	0.73	0.05	-0.26	0.25	•
Psi Blast		2.6e-13	3.4c-10	2.6e-13	3,4e-06	3.46-20	3.4e-20
END		81	. • 76	78	82	121	121
START AA		31	33	32	33	29	6
CHAIN		¥	V	Ą	Ą		
PDB ID		lfbv	1fbv	1g25	1g25	1rmd	1rmd
SEQ ID NO:		32		32	32	32	32

		_		·	· ·											·				
PDB annotation	ANTIBODY, MAD, RING FINGER, 2 ZINC BINUCLEAR CLUSTER, ZINC FINGER, DNA-BINDING PROTEIN		STRUCTURAL PROTEIN TWO REPEATS OF SPECTRIN, ALPHA HELICAL LINKER REGION, 2.2 TANDEM 3-HELIX COLLED-	PROTEIN	BNDOCYTOSIS/EXOCYTOSI S NSEC1; PROTEIN-PROTEIN COMPLEX, MULTI-SUBUNIT	ENDOCYTOSIS/EXOCYTOSI	S SYNAPTOTAGMIN ASSOCIATED 35 KDA	PROTEIN, P35A, THREE HELIX BUNDLE	PROTEIN TRANSPORT HELIX-TURN-HELIX TPR-		TRANSPORT		SIGNALING PROTEIN GTP- BINDING PROTEINS,	COMPLEX, EFFECTORS	5 0	GTP-BINDING PROTEIN IL	SMALL G PROTEIN, RAP2,	GDP, KAS	COMPLEX (GTP- BINDING/EFFECTOR) RAS- [[]	RELATED PROTEIN RABSAŢŲ COMPLEX (GTP-
Compound			ALPHA SPECTRIN; CHAIN: A, B, C;		SYNTAXIN BINDING PROTEIN 1; CHAIN: A; SYNTAXIN 1A; CHAIN: B:	SYNTAXIN-1A; CHAIN: A,	ာ်		VESICULAR TRANSPORT PROTEIN SEC17: CHAIN:	A;			RAS-RELATED PROTEIN RAP-1A; CHAIN: A;	PROTO-UNKOGENE SERINE/THREONINE	PROTEIN KINASE CHAIN: B;	RAP2A; CHAIN: NULL;			RAB-3A; CHAIN: A; RABPHILIN-3A; CHAIN: B;	
SEQ FOLD score			·				÷						123.90			117.80			152.38	
PMF score			0.04		0.27	-0.09			0.04											
Verify score			0.36		-0.03	0.09			0.19											
Psi Blast			2.6e-14		6.5e-18	2.6e-14	•		6.5e-05				1.4e-64			6.8e-58		97.07	0.86-68	
END AA			137		166	120	_		92				161			861	· · · · · · · · · · · · · · · · · · ·	٤	707	1.
START AA			8		7	7			22				25 25			.32	<u> </u>	1		
CHAIN			4		В	Y			∢				∢							
PDB TD			lcun		ldn1	lez3			Iqqe				lcly			1kao		7.1.1	pozi	
SEQ ID NO:			33		33	33			33				34 			34	-	100		

	O	ain, B 2 B 2	O; GMA TON	TOR) IIA; A O	SDZ/D Év	i E	
PDB annotation	FFECTOR) JEFECTOR, SYNAPTIC IIS, RAB AB3A,	IE G PROTI STRAFFICI SLYSIS, RA NSMITTER	TION IN SIGMAT THERASE SIGMAT TENSION IN SIGMAT	BLOOD TONMHIB HROMBIN E, SERINE E), PLASM INDING, 2 IEIN, COM	ULANT AN T, PEPTIDI TIONAL 2 Y, SERINE NHIBITOR	DING PROT N, HEVEIN JA,	SOI ENGLY INCH
PDB	BINDING/BEFECTOR), GPROTEIN, EFFECTOR, RABCDR, 2 SYNAPTIC EXOCYTOSIS, RABPROTEIN, RAB3A,	KABFHILIN HYDROLASE G PROTEIN, VESICULAR TRAFFICKING, GTP HYDROLYSIS, RAB 2 PROTEIN, NEUROTRANSMITTER RELEASE, HYDROLASE	TRANSCRIPTION REGULATION SIGMA70; RNA POLYMERASE SIGMA FACTOR, TRANSCRIPTION REGIT ATTON	COMPLEX (BLOOD COAGULATION/INFIBITOR) AUTOPROTHROMBIN IIA; HYDROLASE, SERINE PROTEINASE), PLASMA CALCIUM BINDING, 2 GLYCOPROTEIN, COMPLEX (BLOOD	ANTI-COAGULANT ANTI- COAGULANT, PEPTIDIC INHIBITORS, CONFORMATIONAL 2 FLEXIBILITY, SERINE PROTEASE INHIBITOR	SUGAR BINDING PROTEIN UDA; LECTIN, HEVEIN DOMAIN, UDA, STIDERA ANTIGEN	יייייייייייייייייייייייייייייייייייייי
pu		÷	SE A : NULL;	HB-PRO-	AIN:		1
Compound	. •	RAB3A; CHAIN: A;	RNA POLYMERASB PRIMARY SIGMA FACTOR; CHAIN: NULL;	ACTIVATED PROTEIN C; CHAIN: C, L; D-PHE-PRO- MAI; CHAIN: P;	HRUSTASIN; CHAIN: NULL;	AGGLUTININ ISOLECTIN VI/AGGLUTININ ISOLECTIN V; CHAIN: A;	
SEQ FOLD score		R 169.98	79.28 R	S0.24 A	HZ	V V	
PMF					-0.17	0.65	
Verify Score					0.03	0.04	
Psi Blast		1.4e-68	5.2e-05	2.6e-08	3.96-11	2.6e-09	
END	·	198	432	233	105	194	3
START AA		29	119	128	2	115	-
CHAIN		A		ц		∢	
PDB CI		3rab	1sig	laut	/xgr	leis	١
SEQ ID NO:		34	40			41	. 17

Table 5

PDB annotation	UDA; LECTIN, HEVEIN DOMAIN, UDA, SUPERANTIGEN	SUGAR BINDING PROTEIN UDA; LECTIN, HEVEIN DOMAIN, UDA, SUPERANTIGEN	SUGAR BINDING PROTEIN CUDA; LECTIN, HEVEIN DOMAIN, UDA, SUPERANTIGEN, SACCHARIDE BINDING	SUGAR BINDING PROTEIN UDA; LECTIN, HEVBIN DOMAIN, UDA, SUPERANTIGEN, SACCHARIDE BINDING	SUGAR BINDING PROTEIN UDA; LECTIN, HEVEIN DOMAIN, UDA, SUPERANTIGEN, SACCHARIDE BINDING	GLYCOPROTEIN TO		COMPLEX (BLOOD U) COAGULATION/INHIBITOR) CHRISTMAS FACTOR; [1] COMPLEX, INHIBITOR, HEMOPHILIA/EGF, BLOOD COAGULATION, 2 PLASMAP SERINE PROTEASE, CALCTUM-BINDING, [1] HYDROLASE, 3 [1] GLYCOPROTEIN
Compound	VI/AGGLUTININ ISOLECTIN V; CHAIN: A;	AGGLUTININ ISOLECTIN VI/AGGLUTININ ISOLECTIN V; CHAIN: A;	AGGLUTININ ISOLECTIN I/AGGLUTININ ISOLECTIN V/ CHAIN: A;	AGGLUTININ ISOLECTIN I/AGGLUTININ ISOLECTIN V/ CHAIN: A;	AGGLUTININ ISOLECTIN V/CHAIN: A;	LAMININ; CHAIN: NULL;	AGGREGATION INFIBITOR, GP ANTAGONIST KISTRIN (NMR, 8 STRUCTURES) IKST 3	FACTOR IXA; CHAIN: C, L,; D-PHE-PRO-ARG; CHAIN: I;
SEQ FOLD score						70.40		59.55
PIMF score		0.41	0.62	0.11	0.03		0.00	
Verify score		-0.10	-0.12	-0.71	-0.05		-0.45	
Psi Blast		1.3e-08	1.3e-10	9.1e-11	1.3e-10	6.5e-22	1.3e-05	3.9e-16
END		169	194	74	169	221	155	148
START AA		. 06	113	9	85	65	. 63	17
CHAIN		Ą	A	¥	Ā			µ
PDB TD		1eis	len2	len2	len2	1klo	1kst	1pfx
SEQ ID NO:		41	41	41	41	41	41	41

Table 5

		_	,4	ص	ــــــ										_					_							
PDB annotation			GLYCOPROTEIN	MEMBRANE COFACTOR .	RECEPTOR, COMPLEMENT	COFACTOR, SHORT	CONSENSUS REPEAT, 2 SCR,	MEASLES VIRUS, GLYCOPROTEIN	GLYCOPROTEIN	MEMBRANE COFACTOR PROTEIN (ACP): VIRIS	RECEPTOR COMPLEMENT	COFACTOR, SHORT	CONSENSUS REPEAT, 2 SCR,	MEASLES VIRUS,	GLYCOPROTEIN	HYDROLASE/HYDROLASE INHIBITOR PROTEIN-	PEPTIDE COMPLEX				COMPLEMENT INHIBITOR (I)	NAR MODILES PROTEIN	STRUCTURE, VACCINIA	VIRUS	COMPLEMENT INHIBITOR H	Z	VIRUS ACCUMA III
Compound	MEMBRANE PROTEIN VITELLINE MEMBRANE OUTER LAYER PROTEIN I 1VMO 3	•	CD46; CHAIN: A, B, C, D, B,	ŗ,					CD46; CHAIN: A, B, C, D, B,	Ĭ,						DES-GLA PACTOR VIIA (HEAVY CHAIN); CHAIN:	H, I; DES-GLA FACTOR	VIIA (LIGHT CHAIN); CHAIN: I. M. COPN)-PHR-	ARG: CHAIN: C. D.	PEPTIDE E-76; CHAIN: X, Y:	COMPLEMENT CONTROL	PROTEIN; CHAIN: A;			COMPLEMENT CONTROL		
SEQ FOLD score																											
PMF	-0.19		-0.08						0.94							0.0 24					0.27				-0.02		
Verify score	0.04		0.31	÷				· .	-0.09							0.14					0.40				0.29		
Psi · Blast	5.2e-38		2.6e-11						1.3e-17							5.1e-12					2.6e-09			-	2.6e-12		
END AA	196		1597						1647							924					1523				9651		
START AA	28		1464						1525							836					1395	·			1467		
CHAIN	Ą		A						٧							H					Ą				٧		
PDB ED	lvmo		1ckl						1ckl							ldva					1e5g				1e5g		
SEQ ID NO:	41		44						4							4					4	•			44		

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PDB annotation	COMPLEMENT INFIBITOR VCP, SP35; COMPLEMENT, NMR, MODULES, PROTEIN STRUCTURE, VACCINIA VIRUS	MATRIX PROTBIN EXTRACELLULAR MATRIX CALCIUM-BINDING, GLYCOPROTBIN, 2 REPEAT, SIGNAL, MULTIGENE FAMILY, DISEASE MUTATION, 3 EGF-LIKE DOMAIN, HUMAN FIBRILLIN-1 FRAGMENT,	MATRIX PROTEIN MATRIX PROTEIN EXTRACELLULAR MATRIX, CALCIUM-BINDING, GLYCOPROTEIN, 2 REPEAT, SIGNAL, MULTIGENE FAMILY, DISEASE MUTATION, 3 EGP-LIKE DOMAIN, HUMAN FIBRILIN-1 FRAGMENT,	BLOOD CLOTTING COMPLEX(SERINE PROTEASE/COFACTOR/LIG AND), BLOOD COAGULATION, 2 SERINE (I) PROTEASE, COMPLEX, CO-C) FACTOR, RECEPTOR FACTOR, RECEPTOR GLA, EGF, COMPLEX GLA, EGF, COMPLEX GLA, EGF, COMPLEX GLA, EGF, COMPLEX GLA, EGF, COMPLEX ALOOD CLOTTING AND), BLOOD CLOTTING	CELL ADHESION PROTEIN ȚŲ EGP-LIKE DOMAIN, CELL ȚŲ
Compound	COMPLEMENT CONTROL PROTEIN; CHAIN: A;	FIBRILLIN; CHAIN: NULL;	FIBRILLIN; CHAIN: NULL;	BLOOD COAGULATION FACTOR VIIA; CHAIN: L; BLOOD COAGULATION FACTOR VIIA; CHAIN: H; SOLUBLE TISSUB FACTOR; CHAIN: T; 5L15; CHAIN: I;	P-SELECTIN; CHAIN: NULL;
SEQ FOLD score			·		
PMF score	0.17	-0.17	-0.20	6.13	-0.20
Verify score	-0.33	0.17	0.35	0.03	0.25
Psi Blast	3.9e-10	1.7e-09	1.7e-09	5.1e-12	5.1e-09
END AA	1633	1127	137	924	143
START AA	1525	1054	73	836	108
CHAIN	∀			ы	
PDB ID	1e5g	lemn	1emn	1 fak	1fsb
SEQ ID NO:	44	4	44	4	44

							- Aleman - A
PDB annotation	ADHESION PROTEIN, TRANSMEMBRANE, 2 GLYCOPROTEIN		GLYCOPROTEIN	SERING PROTEASE FVIIA; FVIIA; BLOOD COAGULATION, SERINE PROTEASE	METAL BINDING PROTEIN BETA SANDWICH, CALCIUM-BINDING PROTEIN, METAL BINDING	MEMBRANE ADHESION SHORT CONSENSUS REPEAT, SUSHI, COMPLEMENT CONTROL PROTEIN, 2 N- GLYCOSYLATION, MULTI- DOMAIN, MEMBRANE	MEMBRANB ADHESION SHORT CONSENSUS REPEAT, SUSHI, COMPLEMENT CONTROL PROTEIN, 2 N- GLYCOSYLATION, MULTI- DOMAIN, MEMBRANE II
Compound		GLYCOPROTEIN FACTOR H, 15TH C.MODULE PAIR (NMR, MINIMIZED AVERAGED 1HFIA 1 STRUCTURE) 1HFIA 1	LAMININ; CHAIN: NULL;	COAGULATION FACTOR VIIA (LIGHT CHAIN); CHAIN: L; COAGULATION FACTOR VIIA (HEAVY TRIPHTIDYL INHIBITOR; CHAIN; CHAIN: H;	LAMININ ALPHA2 CHAIN; CHAIN: A, B, C, D;	HUMAN BETA2- GLYCOPROTEIN I; CHAIN: A;	HUMAN BETA2- GLYCOPROTBIN I; CHAIN: A;
SEQ FOLD score						113.83	
PMF score	·	0.05	-0.18	0.03	-0.12		0.22
Verify score		0.04	0.00	0.14	-0.00		0.16
Psi Blast		2.6e-09	3.4e-10	3.4e-11	7.8e-12	3.9e-19	1.7e-12
END AA		1596	526	924	434	1780	1709
START AA	·	1522	335	841	283	1456	1461
CHAIN ID				1	¥	A	4
PDB ID		Ihfi	1 klo	lqfk	Iqu0	Iqub	1qub
SEQ ID NO:		4	44	4	44	4	4

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PDB annotation	COMPLEMENT INHIBITOR SP35, VCP, VACCINIA VIRUS SP35; COMPLEMENT INHIBITOR, COMPLEMENT MODULE, SCR, SUSHI DOMAIN, 2 MODULE PAIR	COMPLEMENT INHIBITOR SP35, VCP, VACCINIA VIRUS SP35; COMPLEMENT INHIBITOR, COMPLEMENT MODULE, SCR, SUSHI DOMAIN, 2 MODULE PAIR					PROTEIN BINDING ACTIN-U) BINDING PROTEIN,	ILE PROTEIN SFILIN ISOFORN DING PROTEIN, ROLINE ROTEIN, ILE PROTEIN	STRUCTURAL PROTEIN MI
Compound	VACCINIA VIRUS COMPLEMENT CONTROL PROTEIN; CHAIN: NULL;	VACCINIA VIRUS COMPLEMENT CONTROL PROTEIN; CHAIN: NULL;	LECTIN (AGGLUTININ) WHEAT GERM AGGLUTININ (ISOLECTIN 2) 9WGA 3	LECTIN (AGGLUTININ) WHEAT GERM AGGLUTININ (ISOLECTIN 2) 9WGA 3	PROTEIN BINDING PROFILIN I LACF 3 PROTEIN BINDING, PROFILIN LACF 4A	PROTEIN BINDING PROFILIN I 1ACF 3 PROTEIN BINDING, PROFILIN 1ACF 4A	PROFILIN; CHAIN: NULL;	PROFILIN II; CHAIN: A, B, C, D;	PROFILIN II; CHAIN: A, B;
SEQ FOLD score		·				92.69			
PMF score	-0.18	0.01	-0.17	-0.19	1.00		1.00	0.94	1.00
Verify score	0.06	-0.14	0.09	0.09	1.04	·	16:0	0.70	0.75
Psi Blast	1.2e-09	3.9e-13	1.7e-14	5.10-12	8.5e-38	8.5e-38	1.4e-46	3.4e-22	1.2e-39
END	1450	1623	1252	1284	128	129	128	125	128
START AA	0561	1524	1072	1123	14	3	81	16	14
CHAIN	·		Ą	¥				∢	A
PDB ID	lvvc	lvvc	9wga	9wga	lacf	lacf	Icqa	1d1j	12k
SEQ ID NO:	44	44	44	44	45	45	45	45	45

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und PDB annotation	SEVEN-STRANDED INCOMPLETE ANTIPARALLEL UP-AND- DOWN BETA 2 BARREL, ACTIN-BINDING PROTEIN, POLY-L-PROLINE BINDING 3 PROTEIN, PP2 BINDING PROTEIN			3	IN: A, B; ACTIN-BINDING PROTEIN, ACTIN-BINDING PROTEIN, PROPELIN, CYTOSKEL FTON		1	-			-
Compound		PROFILIN; CHAIN: NULL;	PROFILIN; CHAIN: A, B;	ACTIN BINDING PROTEIN PROFILIN 1PNE 3	PROFILIN; CHAIN: A, B;	PROFILIN; CHAIN: A, B;	PROFILIN I; CHAIN: NULL;		SUMO-1; CHAIN: NULL;	UBIQUITIN-LIKE PROTEIN 7, RUB1; CHAIN: A;	1D8 UBIQUITIN; CHAIN: A;
SEQ FOLD score						70.17					
PMF score		0.94	0.99	0.98	1.00		1.00		0.00	1:00	1.00
Verify score		0.15	99'0	0.41	0.54		0.72		0.28	1.02	0.86
Psi Blast		1.4e-21	le-45	1.2e-20	3.4e-38	3.4e-38	3.4e-44	,	6.8e-05	1.2e-23	1e-31
END		. 125	128	125	125	129	126		105	102	105
START AA	•	16	16	16	14	5	17		ο.	31	31
CHAIN ID			A		Ą	A				Ą	A
PDB ID		Ifil		Ipne	1ypr	lypr	3nul	 - -		1bt0	1c3t
SEQ ID NO:	·	45	45	45	45	45	45	ļ	47	47	47

											
PDB annotation	HYDROPHOBIC CORE, PACKING, ROTAMERS, ROC, 2 UBIQUITIN, DE NOVO PROTEIN TRIOTITIN	SIGNALING PROTEIN NEDD8; NEDD-8, UBIQUITIN-LIKE, PROTEOLYSIS, SIGNALING			UBIQUITIN UBIQUITIN, DESIGNED CORE MUTANT	TRANSCRIPTION TUMOR SUPPRESSOR, CANCER,	SANDWICH, 2 TRANSCRIPTION, TRANSCRIPTIONAL	NOT THE REAL PROPERTY.	GENE REGULATION POZ U DOMAIN; PROTEIN-	TRANSCRIPTIONAL 2 REPRESSOR, ZINC-FINGER (I) PROTEIN, X-RAY CRYSTALLOGRAPHY, 3 PROTEIN STRUCTURE, III PROMYELOCYTIC LEUKEMIA, GENB REGIL ATION	
Compound		UBIQUITIN-LIKB PROTEIN NEDD8; CHAIN: A, B, C, D;	UBIQUITIN	CHROMOSOMAL PROTEIN UBIOUTIN 1UBI 3	UBIQUITIN CORB MUTANT 1D7; CHAIN: A;	ELONGIN B; CHAIN: A, D, G, J; ELONGIN C; CHAIN: B H W, MH	F, L L;		PROMYBLOCYTIC LEUKEMIA ZINC FINGER PROTEIN PLZF; CHAIN: A;		OXIDOREDUCTASE(OXYG EN(A)) GALACTOSE OXIDASE (B.C.1.1.3.9) (PH 4.5) 1GOF 3
SEQ FOLD score						·					
PMF score		1.00	1.00	1.00	1.00	06.0	-		0.70		1.00
Verify score		1.10	1.14	1.28	0.94	96.0	•		0.10		0.52
Psi Blast		3.4e-23	6.8e-32	6.8e-34	1.7e-32	5.1e-05			6.8e-22		le-12
AA EB		103	101	105	105	68			193		655
START AA		31	31	31	31	31			02		339
CHAIN		Ą	В		Ą	∢			¥		
EDB ID		1ndd	1tbe	lubi	1ud7	1vcb			1buo		lgof
SEQ ID NO:		47	47	47	47	47			48		48

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PDB annotation		SERINE PROTEASE	INHIBITOR ALPHA-1-	PROTEINASE INHIBITOR,	ALPHA-1-ANTIPROTEINASE;	SERINE PROTEASE	INHIBITIOR, SERPIN,	GLYCOPROTEIN, SIGNAL, 2	POLYMORPHISM,	EMPHYSEMA, DISEASE	SEBINE BROWN SE	NETRITOR AT DEA 1	PROTEINASE INFIBITOR.	ALPHA-1-ANTIPROTEINASE;	SERINE PROTEASE	INHIBITOR, SERPIN,	GLYCOPROTEIN, SIGNAL, 2	POLYMORPHISM,	EMPHYSEMA, DISEASE	MUTATION, ACUIE FRASE	SERPIN AACT SERPIN, SERINE PROTEINASE	INHIBITOR, PARTIAL LOOP	2 INSERTION, LOOP-SHEET	POLYMERIZATION,	EMPHYSEMA, DISEASE 3	MOIAIION, ACOIE PHASE	CONFORMATIONAL	DISEASE	<u> </u>	ur'	real!		COMPLEX (TRANSCRIPTION F	TACTOR/DNA) COMPLEX (TRANSCRIPTION
Compound		ALPHA-1-ANTITRYPSIN;	CHAIN: A;								AT PHA 1 ANTITED VECTAL	CHAIN: 4.							•	, , , , , , , , , , , , , , , , , , , ,	ALPHA-1- ANTICHYMOTRYPSIN;	CHAIN: A;					•		PROTEINASE INHIBITOR	ALFHAI	ANTICHTIMOTRYPSIN		T PROTEIN; CHAIN: A, B;	DNA; CHAIN: C, D;
SEQ FOLD score		300.18												-	-	·							•				·						178.20	
PMF							·	<u>.</u>		. •	9	2								5	90.		-		-		-		90.0					
Verify score						,					0.76)				:				220	7/7								-0.73					
Psi Blast		O									0	·								<	- -								1.3e-07				6.8e-44	
END AA		435									434									127	1								433	_	-		335	
START AA	١	26									61									Ş	3								405				138	
CHAIN		∢									4								,	d	¢								m				∢	
ADA ID		dibi									lalp	:								John	min								Zach				1xbr	
SEQ ID NO:	Ş	49									49									40	}								49				 	

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PDB annotation	FACTORDNA), TRANSCRIPTION FACTOR, 2 DNA-BINDING PROTEIN	COMPLEX (TRANSCRIPTION FACTOR/DNA) COMPLEX (TRANSCRIPTION FACTOR/DNA),	TRANSCRIPTION FACTOR, A DNA-BINDING PROTEIN	CELL ADHESION NEURAL CELL ADHESION	CELL ADHESION NEURAL CELL ADHESION	GROWTH FACTOR/GROWTH FACTOR RECEPTOR FGF, FGFR, IMMUNOGLOBULIN- 1 KF STGNAI	TRANSDUCTION, 2 DIMERIZATION, GROWTH FACTOR/GROWTH FACTOR RECEPTOR	IMMUNE SYSTEM CD32; RECEPTOR, FC, CD32, IMMUNE SYSTEM	COMPLEX (NUCLEOCAPSID. PROTEIN/RNA) NUCLEOCAPSID PROTEIN, COMPLEX (NUCLEOCAPSID) PROTEIN/RNA), 2 STEM-	COMPLEX (NUCLEOCAPSID., PROTEIN/RNA) NUCLEOCAPSID PROTEIN, LA COMPLEX (NUCLEOCAPSID PROTEIN/RNA), 2 STEM- TOOP RNA	
Compound		T PROTEIN; CHAIN: A, B; DNA; CHAIN: C, D;		AXONIN-1; CHAIN: A;	AXONIN-1; CHAIN: A;	FIBROBLAST GROWTH FACTOR 2; CHAIN: A, B; FIBROBLAST GROWTH FACTOR PECTEDIO 1.	CHAIN: C, D;	FC GAMMA RIIB; CHAIN: A;	NUCLEOCAPSID PROTEIN; CHAIN: A; SL3 STEM- LOOP RNA; CHAIN: B;	NUCLEOCAPSID PROTEIN; CHAIN: A; SL3 STEM- LOOP RNA; CHAIN: B;	NUCLEOCAPSID PROTEIN
SEQ FOLD score											
PMF		1.00		90.0	0.37	0.28		0.18	0.27	-0.13	0.12
Verify score		99.0		-0.71	-0.31	-0.49		-0.30	0.07	0.05	0.28
Psi Blast		6.8e-44		1.2e-05	1.3e-05	6.5e-05		5.2e-05	3.4e-17	3.4e-08	6.80-17
END		330		7.1	106	71		86	178	181	178
START AA		140		14	21	17		21	125	151	125
CHAIN		A		A	A	Q		A	¥	ď	
PDB ID	,	1xbr		1cs6	1086	lcvs		2fcb	lalt	lalt	laaf
SEQ ID NO:		20		51	51	51		51	53	53	53

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		1					02/0	
	COMPLEX (NUCLEOCAPSID PROTEIN/DNA) (12-53)NCP7; COMPLEX (NUCLEOCAPSID PROTEIN/DNA), NUCLEIC ACID, 2 RETROVIRUS, VIRUS, MORPHOGENESIS, TANGETERMORPHOGENESIS,	ANTIFREEZE PROTEIN CSPB BETA BARREL, HOMODIMER				NUCLEOCAPSID PROTEIN, NUCLEOCAPSID PROTEIN, HIV-2, RNA RECOGNITION, ZNC BINGED	NUCLEOCAPSID PROTEIN, NUCLEOCAPSID PROTEIN, HTV-2, RNA RECOGNITION, ZINC FINGER	COMPLEX (MHC/VIRAL PEPTIDERECEPTOR) HLA- A2 HEAVY CHAIN; CLASS I MHC, T-CELL RECEPTOR,
HIV-1 NUCLEOCAPSID PROTEIN (MN ISOLATE) (NMR, 20 STRUCTURES) 1AAF 3	DNA (ACGCC); CHAIN: D; NUCLEOCAPSID PROTEIN 7; CHAIN: A;	COLD-SHOCK PROTEIN; CHAIN: A, B;	TRANSCRIPTION REGULATION MAJOR COLD SHOCK PROTEIN (CSPR) 1CSP 3	TRANSCRIPTION REGULATION MAJOR COLD SHOCK PROTEIN	TRANSCRIPTION REGULATION MAJOR COLD SHOCK PROTEIN 7.4 (CSPA (CS 7.4)) OF IMIC 3 (ESCHERICHIA COLI) IMIC 4	NUCLEOCAPSID PROTEIN; CHAIN: NULL;	NUCLEOCAPSID PROTEIN; CHAIN: NULL;	HLA-A 0201; CHAIN: A; BETA-2 MICROGLOBULIN; CHAIN: B; TAX PEPTIDE; CHAIN: C; T CELL
								65.08
	0.15	0.69	0.40	0.87	0.83	0.06	-0.06	
	-0.14	0.29	0.20	0.64	0.21	-0.02	0.26	:
	16-14	1.7e-10	1.7e-10	1.3e-19	le-10	6.8e-05	5.1e-08	3.4e-34
	178	107	107	112	106	159	181	134
	138	40	5	4	36	133	154	21
	∢	4			·			Q
	15j6	1090	Icsp	Icsp	1mjc	1nc8	lnc8	1ao7
	<u> </u>	53	ς 		53	53	23	24

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					Title after trans		
PDB annotation	VIRAL PEPTIDB, 2 COMPLEX (MHC/VIRAL PEPTIDE/RECEPTOR	COMPLEX (MHC/VIRAL PEPTIDERECEPTOR) HLA- A2 HEAVY CHAIN; CLASS I MHC, T-CELL RECEPTOR, VIRAL PEPTIDE, 2 COMPLEX (MHC/VIRAL PEPTIDE/RECEPTOR	IMMUNOGLOBULIN IMMUNOGLOBULIN, ANTIBODY, FAB, ENZYME INHIBITOR, PCR, 2 HOT START	IMMUNOGLOBULIN IMMUNOGLOBULIN, ANTIBODY, FAB, ENZYME INHIBITOR, PCR, 2 HOT START	T CELL RECEPTOR TCR; T CELL RECEPTOR, MHC CLASS I, HUMAN IMMUNODEFICIENCY VIRUS, 2 MOLECULAR RECOGNITION	T CELL RECEPTOR TCR; T CELL RECEPTOR, MHC (I) CLASS I, HUMAN INMUNODERICIENCY II VIRUS, 2 MOLECULAR SECOGNITION	COMPLEX (MHCVIRAL PEPTIDERECEPTOR) HLA PA A2 HBAVY CHAIN; IU COMPLEX (MHCVIRAL PEPTIDERECEPTOR) III
Compound	RECEPTOR ALPHA; CHAIN: D; T CELL RECEPTOR BETA; CHAIN: E;	HLA-A 0201; CHAIN: A; BETA-2 MICROGLOBULIN; CHAIN: B; TAX PEPTIDE; CHAIN: C; T CELL RECEPTOR ALPHA; CHAIN: D; T CELL RECEPTOR BETA; CHAIN: B;	TP7 FAB: CHAIN: L, H;	TP7 FAB; CHAIN: L, H;	T CELL RECEPTOR V- ALPHA DOMAIN; CHAIN: A, B;	T CELL RECEPTOR V. ALPHA DOMAIN; CHAIN: A, B;	HLA-A 0201; CHAIN: A; BETA-2 MICROGLOBULIN; CHAIN: B; TAX PEPTIDE; CHAIN: C; T CELL RECEPTOR ALPHA;
SEQ FOLD score		54.73		51.86	67.06		·
PMF score			0.30	·		0.92	1.00
Verify			0.24			0.30	0.26
Psi Blast	,	5.1e-28	6.8e-35	6.8e-35	1.7e-38	1.7e-38	1.7e-37
END		203	148	204	130	142	196
START AA		21		19	20	20	20
CHAIN		В	1	1	Y	V	D
PDB CI		1ao7	layl	layl	1688	1588	1bd2
SEQ ID NO:	·	54	54	54	54	54	54

				# 'P # ''		beet "12 seems gewate screen
PDB annotation		COMPLEX (MHC/VIRAL PEPTIDERECEPTOR) HLA A2 HEAVY CHAIN; COMPLEX (MHC/VIRAL PEPTIDERECEPTOR)	IMMUNE SYSTEM IMMUNOGLOBULN, IMMUNORECEPTOR, IMMUNE SYSTEM	IMMUNE SYSTEM MHC I- AK; MHC I-AK; T-CELL RECEPTOR, MHC CLASS II, D10, I-AK	IMMUNE SYSTEM FAB-IBP COMPLEX CRYSTAL STRUCTURE 2.7A RESOLUTION BINDING 2 OUTSIDE THE ANTIGEN COMBINING SITE SUPERANTIGEN FAB VH3 3 SPECIFICITY	COMPLEX (HIV ENVELOPE PROTEIN/CD4/FAB) COMPLEX (HIV ENVELOPE PROTEIN/CD4/FAB), HIV-1
Compound	CHAIN: D; T CELL RECEPTOR BETA; CHAIN: E;	HLA-A 0201; CHAIN: A; BETA-2-MICROGLOBULIN; CHAIN: B; TAX PEPTIDE; CHAIN: C; T CELL RECEPTOR ALPHA; CHAIN: D; T CELL RECEPTOR BETA; CHAIN: B;	ALPHA-BETA T CELL. RECEPTOR (TCR) (D10); CHAIN: A;	T-CELL RECEPTOR D10 (ALPHA CHAIN); CHAIN: A, B; T-CELL RECEPTOR D10 (BETA CHAIN); CHAIN: B, F; MHC I-AK A CHAIN: CHAIN; CHAIN: C, G; MHC I-AK B CHAIN: C, G; MHC I-AK B CHAIN: D, H; CONALIBUMIN PEPTIDE;	GENERALY, CHAIN: A, C, IGM RF 2A2; CHAIN: A, C, E; IGM RF 2A2; CHAIN: B, D, F; IMMUNOGLOBULIN G BINDING PROTEIN A; CHAIN: G, H;	ENVELOPB PROTEIN GP120; CHAIN: G; CD4; CHAIN: C; ANTIBODY 17B; CHAIN: L, H;
SEQ FOLD score		70.90				50.38
PMF score			0.88	1.00	0.52	
Verify score			0.13	0.51	0.12	
Psi Blast		1.7e-37	3.4e-40	6.86-39	6.8e-39	6.8e-33
END		204	129	129	165	204
START AA		20	61	21	19	19
CHAIN		Ω	Ą	Y	α .	1
PDB TD		1bd2	Ibwm	1 d9k	1dee	lgc1
SEQ ID NO:		54	54	54	54	54

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PDB annotation	GP120, T-CELL SURFACE GLYCOPROTEIN CD4, 3 ANTIGEN-BINDING FRAGMENT OF HUMAN IMMUNOGLOBULIN 17B, 4 GLYCOSYLATED PROTEIN	IMMUNOGLOBULIN INTACT IMMUNOGLOBULIN V REGION CREGION, IMMUNOGLOBULIN	COMPLEX (ANTIBODY/ANTIGEN) CYTOKINE RECEPTOR, COMPLEX (ANTIBODY/ANTIGEN), 2 TRANSMEMBRANE, GLYCOPROTEIN	COMPLEX (IMMUNOGLOBULINIRECEP TOR) TCR VAPLHA VBETA DOMAIN; T-CELL RECEPTOR, STRAND SWITCH, FAB, ANTICLONOTYPIC, 2 (IMMUNOGLOBULINIRECEP)	COMPLEX . (IMMUNOGLOBULIN/RECEP TOR) TCR VAPLHA VBETA (DOMAIN; T-CELL RECEPTOR, STRAND SWITCH, PAB, ANTICLONOTYPIC, 2 (IMMUNOGLOBULIN/RECEP TOR)	INMUNE SYSTEM HUMAN TCR/PEPTIDE/MHC COMPLEX, HLA-A2, HTLV-1, TAX, TCR, T 2 CELL
Compound		IGG2A INTACT ANTIBODY - MAB231; CHAIN: A, B, C, D	ANTIBODY A6: CHAIN: L, H; INTERFERON-GAMMA RECEPTOR ALPHA CHAIN; CHAIN: I;	KB5-C20 T-CELL ANTIGEN RECEPTOR; CHAIN: A, B; ANTIBODY DESIRE-1; CHAIN: L, H;	KBS-C20 T-CELL ANTIGEN RECEPTOR; CHAIN: A, B; ANTIBODY DESIRE-1; CHAIN: L, H;	MHC CLASS I HLA-A; CHAIN: A; BETA-2 MICROGLOBULIN; CHAIN: B; TAX PEPTIDE P6A;
SEQ FOLD score	·	·	50.28	62.17		71.85
PMF		0.45			0.11	
Verify score		0.01			4 20	
Psi Blast		3.4e-35	3.46-27	1.7e-40	1.7e-40	1.7c-36
END AA		168	192	134	147	204
START AA		19	61	20	20	21
CHAIN	·	В	1	· V	∀	D
PDB ID		ligt	ljrh	1kb5	1kb5	lqrn
SEQ ID NO:		54	54	54	54	54

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PDB annotation	RECEPTOR, IMMUNE SYSTEM	IMMUNE SYSTEM HUMAN TCR/PEPTIDE/MHC COMPLEX, HLA-A2, HTLV-1,	TAX, TCR, T 2 CELL RECEPTOR, IMMUNE SYSTEM	COMPLEX (COAT PROTEIN/IMMUNOGLOBULI N) POLYPROTEIN, COAT PROTEIN CORR PROTEIN	RNA-DIRECTED RNA 2 POLYMERASE, HYDROLASE, THIOL	MYRISTYLATION, 3 COMPLEX (COAT PROTEIN/IMMUNOGLOBULI N)	RECEPTOR TCR; T-CELL, RECEPTOR, TRANSMEMBRANE, GLYCOPROTEIN, SIGNAL			AMINOPEPTIDASE AMINOPEPTIDASE, PROLÎNE ÎMINOPEPTIDASE, SERINE PROTEASE, 2
Compound	CHAIN: C; HMAN T-CELL RECEPTOR; CHAIN: D; HLA-A 0201; CHAIN: B;	MHC CLASS I HLA-A; CHAIN: A; BETA-2 MICROGLOBULIN: CHAIN:	B; TAX PEPTIDE P6A; CHAIN: C; HMAN T-CELL RECEPTOR; CHAIN: D; HLA-A 0201; CHAIN: E;	HUMAN RHINOVIRUS 14 COAT PROTEIN; CHAIN: 1, 2, 3, 4; FAB 17-IA; CHAIN:	: :		ALPHA, BETA T-CELL RECEPTOR CHAIN: A, B;	IMMUNOGLOBULIN FAB 2FB4 4	IMMUNOGLOBULIN FAB FRAGMENT FROM HUMAN IMMUNOGLOBULIN IGG1 (LAMBDA, HIL.) 8FAB 3	PROLINB IMINOPEPTIDASE; CHAIN: A, B;
SEQ FOLD score				51.58			55.23		50.44	
PMF		1.00						0.55		0.30
Verify		0.38						-0.12		0.10
Psi Blast		1.7e-36		1.7e-31			1.3e-19	6.8e-36	3.4e-32	5.1e-05
END		196		131			204	163	203	205
START		22		19		-	21	19	19	107
CHAIN		Ω		ы			¥	H	В	Α .
PDB CD		Igm		lrvf			Itcr	2fb4	8fab	lazw
SEQ ID		54		54			54	54	54	56

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PDB annotation	XANTHOMONAS CAMPESTRIS	HALOPEROXIDASE HALOPEROXIDASE A2,	CHLOROPEROXIDASE A2; HALOPEROXIDASE,	OXIDOREDUCTASE,	PEROXIDASE, ALPHA/BETA	2 HYDROLASE FOLD,	LIVER OF THE PRING	HYDROLASE DITLE,	DEGRADATION	LIPASE ESTERASE,	SUBSTRATE/PRODUCT-	BOUND ICLE 9	HYDROLASE ALPHA/BETA	HYDROLASE FOLD	HYDROLASE ALPHA/BETA	HYDROLASE FOLD	HYDROLASE BILE SALT	ACTIVATED LIPASE,	ESTERASE, CATALYTIC	DOMAIN	SERINE HYDROLASE	LIPASE	SERINE HYDROLASE	SERINE HYDROLASE,	DEGRADATION OF	BREFELDIN A, ALPHA/BETA	2 HYDROLASE FAMILY	SERINE HYDROLASE	DEGRADATION OF	BREFFI DIN A ALPHA/BETA	2 HYDROLASE FAMILY	meia, Utaur
Compound		BROMOPEROXIDASE A2; CHAIN: NULL;					2 HYDDOXY 6 OYO 6	PHRNYI HEXA-2 4.	DIENOATE CHAIN: A;	CHOLESTEROL	ESTERASE; 1CLE 4 CHAIN:	A, B; 1CLE 5	SERINE HYDROLASE;	CHAIN: A;	SERINE HYDROLASE;	CHAIN: A;	BILE SALT ACTIVATED	LIPASE; CHAIN: A;			LIPASE; CHAIN: A, B;	,	BREFELDIN A ESTERASE;	CHAIN: A, B;				BREFELDIN A ESTERASE;	CHAIN: A, B;			HYDROLASE(CARBOXYLI C ESTERASE) LIPASE
SEQ FOLD score										63.65																		87.04				
PMF score		0.15					0.04	5					0.36		9.1		-0.14			100	0.21		0.55									0.07
Verify score		0.17					0.21	1					-0.18		0.14		0.05		•		-0.35		-0.16									0.18
Psi Blast		0.00051					5 18-07	; ;		8.5e-57			8.5e-13		1.7e-39		5.1e-15				5.1e-06		1.4e-25					1.4e-25				0.00026
END		215					216			515			509		345		513			900	299		321	•				403				246
START AA		08					195	}					411		73		415				55		65					×				. 201
CHAIN							\ 			Ą			∢		∢		∀				∢		Ą					∢				В
PDB ID		lbrt					1c4x			Icle			levq		levq		1f6w				#[1		1jkm				,	1.jkm				11pb
SEQ ID NO:		95					56			56			26	ì	96		26			72	90		56					ج و				26

Table 5

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PDB annotation			HYDROLASE MACHE; HYDROLASE, SERINE ESTERASE, ACETYLCHOLINESTERASE, TETRAMER, 2 HYDROLASE FOLD, GLYCOSYLATED PROTEIN	HYDROLASE PNB ESTERASE; ALPHA-BETA HYDROLASE DIRECTED EVOLUTION	HYDROLASE ALPHA BETA HYDROLASE FOLD, PROLING, PROLYL AMINOPEPTIDASE, 2 SERRATIA,		
Compound	(E.C.3.1.1.3) COMPLEXED WITH COLIPASE AND INHIBITED 11.PB 3 BY UNDECANE PHOSPHONATE METHYL ESTER (TWO CONFORMATIONS) 11.PB 4	HYDROLASE LIPASE (E.C.3.1.1.3) (TRIACYLGLYCEROL LIPASE) COMPLEXED WITH 11.PP 3 HEXADECANESULFONAT E 11.PP 4 11.PP 71	ACETYLCHOLINESTERAS E; CHAIN: A, B, C, D;	PARA-NITROBENZYL ESTERASE; CHAIN: A;	PROLYL AMINOPEPTIDASE; CHAIN: A;	HYDROLASE(CARBOXYLI C ESTERASE) LIPASE (B.C.3.1.1.3) TRIACYLGLYCEROL HYDROLASE 1THG 3	HYDROLASE(CARBOXYLI CESTERASE) LIPASE (E.C.3.1.1.3)
SEQ FOLD score		64.96	62.20	66.32		·	75.37
PMF score					0.40	0.24	
Verify score					-0.31	-0.37	
Psi Blast		5.1e-54	6.8e-87	1.7e-67	1.2e-05	7.8e-22	1.7e-55
END AA			515	511	205	197	497
START AA		T	1	23	\$9	19	1
CHAIN			A	· •	Y		
PDB CD		llpp	Imaa	1 qe 3	lqt	1thg	1thg
SEQ ID NO:		S6	56	. 95	56	56	56

					,		-17**		1.12.17.2		n
PDB annotation		HYDROLASE BILE SALT ACTIVATED LIPASE, BILE SALT STIMULATED HYDROLASE, SERINE ESTERASE, LIPASE	COMPLEX (TRANSCRIPTION PACTOR/DNA) COMPLEX (TRANSCRIPTION PACTOR/DNA), TRANSCRIPTION PACTOR, 2 DNA-BINDING PROTEIN		RIBOSOMAL PROTEIN ALZHEIMER DISEASE, RIBOSOMAL PROTEIN S6, OLIGOMERIZATION	RIBOSOMAL PROTEIN ALZHEIMER DISBASE, RIBOSOMAL PROTEIN S6, OLIGOMERIZATION		SIGNAL TRANSDUCTION FULG, DHR3 DOMAIN; SIGNAL TRANSDUCTION, SH3 DOMAIN, REPEAT	OXIDOREDUCTASE BETA- L FINGER	MEMBRANE PROTEIN/OXIDOREDUCTAS E BETA-FINGER, HETERODIMER	PEPTIDE RECOGNITION
Compound	TRIACYLGLYCEROL HYDROLASE 1THG 3	CHOLESTEROL BSTERASE; CHAIN: NULL;	T PROTEIN; CHAIN: A, B; DNA; CHAIN: C, D;		RIBOSOMAL PROTBIN S6; CHAIN: A, B;	RIBOSOMAL PROTEIN S6; CHAIN: A;		HUMAN DISCS LARGE PROTEIN; CHAIN: NULL;	NEURONAL NITRIC OXIDE SYNTHASE (RESIDUES 1-130); CHAIN: A;	ALPHA-1 SYNTROPHIN (RESIDUES 77-171); CHAIN: A; NEURONAL NITRIC OXIDE SYNTHASE (RESIDUES 1-130); CHAIN: B;	POSTSYNAPTIC DENSITY
SEQ FOLD score		58.21	161.28					·		·	
PMF score					0.80	0.92		0.83	0.70	0.55	0.55
Verify score				•	0.35	0.52		0.48	0.47	0.62	0.49
Psi Blast		5.1e-76	5.1e-84		6.8e-22	1.7e-19		8.4e-17	1.2e-17	1.20-17	3.6e-18
END AA		514	306		66 .	95		255	270	248	249
START AA		16	66		1	I		153	157	153	157
CHAIN			V		Ą	A		•	4	₹	A
PDB ID		2bce	lxbr.		lcqm	Iqjh		1pdr	lqau 	Iqav	1glc
SEQ TD NO:		56	58		92	92		99	99	99	99

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PDB annotation	PSD-95; PDZ DOMAIN, NEURONAL NITRIC OXIDE SYNTHASE, NMDA RECEPTOR 2 BINDING		COMPLEX (BLOOD COAGULATION/INHIBITOR) AUTOPROTHROMBIN IIA:	HYDROLASE, SERINE	PROTEINASE), PLASMA CALCIUM BINDING, 2	GLYCOPROTEIN, COMPLEX	COAGULATION/INHIBITOR)	HYDROLASE INHIBITOR ALL-BETA STRUCTURE,	HYDROL ASE INHIBITOR	ALL-BETA STRUCTURE,	HYDROLASE INHIBITOR	PLANT PROTEIN TWO	HOMOLOGOUS HEVEIN- LIKE DOMAINS	1	HOMOLOGOUS HEVEIN- LIKE DOMAINS	PLANT PROTEIN TWO	HOMOLOGOUS HEVEIN-	PLANT PROTEIN TWO	HOMOLOGOUS HEVEIN-	PLANT PROTEIN TWO	HOMOLOGOUS HEVEIN- LIKE DOMAINS	SUGAR BINDING PROTEIN	
Compound	PROTEIN 95; CHAIN: A;		ACTIVATED PROTEIN C; CHAIN: C, L; D-PHE-PRO- MAI: CHAIN: P:					BOWMAN-BIRK TRYPSIN INHIBITOR; CHAIN: A	TOTAL STATE OF THE	BOWMAN-BIKK IKYPSIN INHIBITOR; CHAIN: A		AGGLUTININ ISOLECTIN	VI; CHAIN: A	AGGLUTININ ISOLECTIN	VI; CHAIN: A	AGGLUTININ ISOLECTIN	VI; CHAIN: A	AGGLUTININ ISOLECTIN	VI; CHAIN: A	AGGLUTININ ISOLECTIN	VI; CHAIN: A	AGGLUTININ ISOLECTIN	ISOLECTIN V; CHAIN: A;
SEQ FOLD score							,																
PMF score			-0.17					-0.07	9.0	71.0		0.30		0.48		-0.18	-	0.35		0.11		-0.17	
Verify score			0.15					0.10	760	0.84		0.80		2.11		1.23		0.75		0.22	<u> </u>	0.98	
Psi Blast			8.4e-09		<u> </u>			1.26-22	00.00	4.86-20		3.6e-18		4.8e-07		1.1e-18		4.8e-17		1.2e-12		6e-20	
END AA			133					151	106	071		110		30		87		134		146		87	
START AA			40					35	,	4		10		1	•	3		57		70		4	
CHAIN	·		7					¥		<		V		٧		Ą		4		Ą	•	A	
PDB UD			laut					lc2a	5	1028		lehd		lehd		1ehd		lehd		1ehd		leis	
SEQ ID NO:			69					.69	9	8		69		69		69		69		69		69	

Table 5

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tation	SIGNALLING PROTEIN BINDING PROTEIN, CYTOKINE, SIGNALLING PROTEIN	PTOR PTOR, FOR			MEMBRANE ADHESION SHORT CONSENSUS		COMPLEMENT CONTROL PROTEIN 2 N.	GLYCOSYLATION, MULTI-	RANE	SE	INHIBITOR FACTOR XA INHIBITOR: ANTISTASIN.	CRYSTAL STRUCTURE	FACTOR XA INHIBITOR, 2	SE	INHIBITOR, THROMBOSIS	SEKINE PROTEASE INHIBITOR FACTOR XA	INHIBITOR; ANTISTASIN,	CRYSTAL STRUCTURE,	FACTOR AN INFIBITOR, 2	SERINE PROTEASE INHIRITOR, THROMBOSIS	SE	INHIBITOR FACTOR XA	INHIBITOR; ANTISTASIN	CRYSTAL STRUCTURE, FACTOR XA INHIBITOR	
PDB annotation	SIGNALLING PROTEIN BINDING PROTEIN, CYTOKINE, SIGNALLII PROTEIN	HORMONE RECEPTOR, HORMONE RECEPTOR, INSULIN RECEPTOR FAMILY	GLYCOPROTEIN GLYCOPROTEIN	GLYCOPROTEIN GLYCOPROTEIN	MEMBRANE ADHES SHORT CONSENSUS	SUSHI,	MENT 2 N.	LATIC	DOMAIN, MEMBRANE ADHESION	SERINE PROTEASE	R FAC	STRU	XA INE	SERINE PROTEASE	K, IH	SEKUNE PROTEASE INHIBITOR FACTOI	R; ANJ	STRU	AA LINE	SERINE PROTEASE INHIRITOR, THROM	SERINE PROTEASE	R FAC	R; AN	XA INT	
2	SIGNALL BINDING CYTOKIN PROTBIN	HORMON HORMON INSULIN FAMILY	YCOPI YCOPI	YCOP	EMBRA FORT C	REPEAT, SUSHI	COMPLEMENT	YCOS	DOMAIN, MADHESION	RINE P	HIBITIC	YSTAI	CTOR	RINEP	HIBIT	HIBITO H	HIBITC	VSTA	CICK	RINE	RENE	HIBITC	HIBIT	SYSTAI	
	ER CY BE	H H A					<u>ဗ</u>	<u>ਂ</u> ਲ	<u> </u>	땅	<u> </u>	5	E	8		3 Z	<u>Z</u>	មីផ	L.	2 8		<u>z</u>	<u>z</u>	<u> </u>	
: 	R;	OWTH OR 1;	LAMININ; CHAIN: NULL;	LAMININ; CHAIN: NULL;	HUMAN BETA2- GLYCOPROTEIN I: CHAIN:					Ä											Ä				
Compound	TUMOR NECROSIS FACTOR RECEPTOR; CHAIN: A, B;	INSULIN-LIKE GROWTH PACTOR RECEPTOR 1; CHAIN: A;	HAIN:	HAIN	TED I					ANTISTASIN; CHAIN:						ANTISTASIN; CHAIN: NIII I :					ANTISTASIN; CHAIN				
آن ک	TUMOR NEC FACTOR REC CHAIN: A, B;	INSULIN-LI PACTOR RE CHAIN: A;	ININ; (ININ; (HUMAN BETA2					ISTASI	ä					ISTASI I	ì				ISTAS	ij		٠.	
	PAC CHA	PAC	LAM	LAM	HUM	Ą;				A	NULL;				-	ANTIS					AM	NGLL;			
SEQ FOLD score													-			~1									
SEQ	54.40			75.71						_						58.52					1				
PMF		-0.19	-0.03		-0.20			٠		-0.17										,	-0.05	}		·	
Verify		0.60	0.84		0.54					0.40											0.02				
Psi	6e-14	2.4e-22	2.4e-23	2.4e-23	4.8e-24			-		6e-17		_				9.6e-19					9.6e-19	}			
END	155	149	143	145	151					112						124					149	}			
START	2	4	3	4	4					13						23					30	3		•	
CHAIN		A			A											:		·							
PDB	 	ligi	lklo	Iklo	1qub					Iskz						Iskz					1 chr	7			
SEQ ID																6									
S	8	8 .	8	8	8					8						8					8	خ ـــــــ			_

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PDB annotation	SERINE PROTEASE INHIBITOR, THROMBOSIS		l-		THE LANGE OF THE PARTY OF THE P	IKANSPOKI PROTEIN SERINE-RICH RNA POLYMERASE I SUPPRESSOR PROTEIN; ARM REPEAT	SIGNALING PROTEIN PEROXISMORE RECEPTOR 1, PTS1-BP, PEROXIN-5, PTS1 PROTEIN-PEPTIDE COMPLEX, TETRATRICOPEPTIDE REPEAT, TPR, 2 HELICAL		CELL ADHESION NEURAL CELL ADHESION	GROWTH FACTOR/GROWTH FACTOR RECEPTOR FGF, FGFR, IMMUNOGLOBULIN- LIKE, SIGNAL TRANSDUCTION, 2 DIMERIZATION, GROWTH FACTOR/GROWTH FACTOR
Compound		MEMBRANE PROTEIN VITELLINE MEMBRANE OUTER LAYER PROTEIN I IVMO 3	LECTIN (AGGLUTININ) WHEAT GERM AGGLUTININ (ISOLECTIN 2) 9WGA 3	LECTIN (AGGLUTININ) WHEAT GERM AGGLUTININ (ISOLECTIN 2) 9WGA 3	TATAL TATAL STREET	KARY OF HERIN ALCHA; CHAIN: A, B; MYC PROTO- ONCOGENE PROTEIN; CHAIN: C, D, B, F;	PEROXISOMAL TARGETING SIGNAL 1 RECEPTOR; CHAIN: A, B; PTS1-CONTAINING PEPTIDE; CHAIN: C, D;		AXONIN-1; CHAIN: A;	FIBROBLAST GROWTH FACTOR 2; CHAIN: A, B; FIBROBLAST GROWTH FACTOR RECEPTOR 1; CHAIN: C, D;
SEQ FOLD score				97.03						
PMF		-0.18	-0.11		0.00	600	0.96		99.0	0.90
Verify		0.46	0.56		80	800	-0.23		0.22	-0.09
Psi Blast		4.8e-28	3.6e-25	3.6e-25	70000	0.00024	0.00024		6e-15	1.2e-13
END AA		147	151	151	272	<u></u>	574		204	202
START		2	2	2	5		371		34	34
CHAIN		Y	¥	Ą	_	€ .	Ψ .		¥	ر ن
PDB ED		lvmo	9wga	9wga	1004	Tee4	1fch		1cs6	Icvs
SEQ ID NO:		. 69	69	69	22	7/			83	

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PDB annotation	RECEPTOR	CELL ADHESION NCAM; NCAM, IMMUNOGLOBULIN	FOLD, GLYCOPROTEIN	GROWTH FACTOR/GROWTH	FACTOR RECEPTOR FOFZ;	GOT TRE DOMAINS	BELONGING TO THE I-SET 2	SUBGROUP WITHIN IG-LIKE	GROWTH FACTOR/GROWTH	PACTOR RECEPTOR FGF2:	FGFR2; IMMUNOGLOBULIN	(IG)LIKE DOMAINS	BELONGING TO THE I-SET 2	SUBGROUP WITHIN IG-LIKE DOMAINS, R-TREFOIL FOLD	IMMUNE SYSTEM FC-	EPSILON RI-ALPHA;	IMMUNOGLOBULIN FOLD,	GLYCOPROTEIN,	RECEPTOR, IGE-BINDING 2	PROTEIN	IMMUNE SYSTEM HIGH	AFFINITY IGE-FC	RECEPTOR; FC(EPSILON)	IGE-FC; IMMUNOGLOBULIN	FOLD, GLYCOPROTEIN,	RECEPTOR, IGE-BINDING 2	PROTEIN, IGE ANTIBODY,	IGE-FC	MEMBRANE PROTEIN CD32:	FC RECEPTOR.	IMMUNOGLOULIN,	LEUKOCYTE, CD32	RECEPTOR, FC, CD32,
Compound		NEURAL CELL ADHESION MOLECULE; CHAIN: A, B,	C, D;	FIBROBLAST GROWTH	PACIOR Z; CHAIN: A, B, C,	FACTOR RECEPTOR 2.	CHAIN: E. F. G. H:		FIREORI AST GROWTH	FACTOR 2: CHAIN: A. B. C.	D; FIBROBLAST GROWTH	FACTOR RECEPTOR 2;	CHAIN: B, F, G, H;		HIGH AFFINITY	IMMUNOGLOBULIN	EPSILON RECEPTOR	CHAIN: A;	•		HIGH AFFINITY	IMMUNOGLOBULIN	EPSILON RECEPTOR	CHAIN: A; IG EPSILON	CHAIN C REGION; CHAIN:	В, D;			FC RECEPTOR FC(GAMMA)RIIA: CHAIN:	A:			FC GAMMA RIB; CHAIN: A;
SEQ FOLD score																	•				·												
PMF score		0.92		0.31		•		. *	0.65	}					0.34		,				12.0								0.29				0.71
Verify score		0.34		0.12		,			0.12						0.04						-0.14		•						0.0 40.				0.33
Psi Blast		1.2e-15		1.1e-15					3.6e-13						3.6e-13						6e-14			•	•			,	4.86-15				3.6e-15
END AA		202		707					203	}					193						202							18	503			1	202
START AA		34		28					35						34						34							,	¥.				34
CHAIN		A		田					Ð) 				_ .	A		_				Ą								∢	-			¥
PDB ID		lepf		lev2					lev2						1229						1f6a								licg				2fcb
SEQ ID NO:		83		83					83	1					83						83								52				83

					Direction of			
PDB annotation	IMMUNE SYSTEM CELL ADHESION NCAM DOMAIN 1; CELL ADHESION, GLYCOPROTEIN, HEPARIN- BINDING, GPI-ANCHOR, 2 NEURAL ADHESION MOLECULE, IMMUNOGLOBULIN FOLD,	SIGNAL	TRANSPORT PROTEIN SERINE-RICH RNA POLYMERASE I SUPPRESSOR PROTEIN; ARM REPEAT	TRANSPORT PROTEIN SERINE-RICH RNA POL YMERASE I SUPPRESSOR PROTEIN; ARM REPEAT	ENDOCYTOSIS/EXOCYTOSI S SYNAPTOTAGMIN ASSOCIATED 35 KDA PROTEIN, P354, THREE	ARMADILLO REPEAT ARMADILLO REPEAT, BETA-CATENIN, CYTOSKELETON	ARMADILLO REPEAT, ARMADILLO REPEAT, BETA-CATENIN, CYTOSKELETON	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING
Compound	NEURAL CELL ADHESION MOLECULE; CHAIN: NULL;		KARYOPHERIN ALPHA; CHAIN: A, B; MYC PROTO- ONCOGENE PROTEIN; CHAIN: C, D, E, F;	KARYOPHERIN ALPHA; CHAIN: A, B; MYC PROTO- ONCOGENE PROTEIN; CHAIN: C, D, E, F;	SYNTAXIN-1A; CHAIN: A, B, C;	BETA-CATENIN; CHAIN: NULL;	BETA-CATENIN; CHAIN: NULL;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE
SEQ FOLD score								·
PMF	0.94		0.68	0.80	-0.14	0.18	0.93	0.59
Verify	0.49		0.35	0.16	0.01	0.20	0.15	-0.27
Psi Blast	2.46-14		4.8e-09	2.4e-05	3.6e-15	2.4e-09	1.2e-09	6.8e-28
END AA	202		225	311	867	532	440	232
START AA	104		11	3 2	737	108	7	156
CHAIN			∢	∢	¥			A
EDB EDB	2пст		1664	1664	lez3	3bct	3bct	lalh
SEQ ID NO:	83		85	85	85	85	85	86

Table 5

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PDB annotation	PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTBIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC ·
Compound	BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B,	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B,	QGSR ZINC FINGER
SEQ FOLD score		·						
PMF		0.1	-0.06	0.53	0.00	0.19	0.21	0.63
Verlfy		-0.41	0.10	0.02	-0.37	-0.21	-0.44	0.11
Pst Blast		3.4e-27	8.5e-27	6.8e-28	8.5e-29	3.4e-25	1.2e-23	1e-29
END AA		260	365	391	733	785	814	870
START		180	291	313	651	707	739	789
CHAIN		A	¥	¥ ·	V	4	A	Ą
PDB ID		lalh	laih	lalh	laih	lalh	lalh	lalh
SEQ ID NO:			98	98	98	98	86	98

			•			
					PLIZIBUE	. Dleer
PDB annotation	FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN		
Compound	PEPTIDE; CHAIN: A; DUPLEX OLIGONUCI EOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER. PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	TRANSCRIPTION REGULATION YBAST TRANSCRIPTION FACTOR ADRI (RESIDUBS 102 - 130) 1ARD 3 (AMINO TERMINAL ZINC FINGER DOMAIN) (NMR, 10 STRUCTURES) 1ARD 4 (ADR1B) 1ARD 5	DNA-BINDING PROTEIN HUMAN ENHANCER- BINDING PROTEIN MBP-1 MUTANT WITH CYS 11 1BBO 3 REPLACED BY ABU (C11ABU) (NMR, 60 STRUCTURES) 1BBO 4
SEQ FOLD score						
PMF score		0.48	0.43	0.31	00:00	0.19
Verify score		-0.05	-0.17	-0.09	69'0-	-0.31
Psi Blast		1.4e-28	7.26-14	5.1e-27	6.8e-06	3.4e-10
END AA		668	928	946	341	897
START		818	846	875	313	848 8
CHAIN		A	V	A		
PDB U		laih	laih	laih	lard	1bbo
SEQ ID NO:		98	86	98	98	98

Table 5

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PDB annotation	COMPLEX (ZINC FINGER/DNA) ZINC FINGER,	PROTEIN-DNA	DESIGN 2 CRYSTAI	STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC FINGER, FINGER,	PROTEIN-DNA	INTERACTION, PROTEIN	DESIGN, 2 CRYSTAL	(ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER,	PROTEIN-DNA	INTERACTION, PROTEIN	DESIGN, 2 CRYSTAL	STRUCTURE, COMPLEX	(EIIVO I IIVO IIIVO IIIVO)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER,	PROTEIN-DNA NATER ACTION PROTEIN	DESIGN 2 CRYSTAL	STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX-(ZINC FINGER)	PROTEIN-DNA	INTERACTION, PROTEIN	DESIGN, 2 CRYSTAL	STRUCTURE, COMPLEX	(ZINC HINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER	PROTEIN-DNA INTERACTION PROTEIN	
Compound	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;				DNA; CHAIN: A; B, D, E; CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;		-		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;	•				DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;	•			DNA; CHAIN: A, B, D, B;	PROTEIN: CHAIN: C. F. G.					DNA; CHAIN: A, B, D, B; CONSENSUS ZINC FINGER	PROTEÍN; CHAIN: C, F, G,	
SEQ FOLD score																														
PMF score	0.81					1.00					0.95						0.03					0.70						0.35		
Verify score	-0.13		,		1	10.0					-0.16			-			-0.70					-0.22						-0.38		
Psi Blast	6.8e-46					8.5e-46	٠.				1.7e-43						3.46-43		_			5.1e-46		· .				1e-46		
END	232	_				260					287			,			337					391						733	``	
START AA	155					179					207		:				263					312						650.		
CHAIN	၁					ວ					၁						ပ					၁				,		ပ		
PDB CD	1mey					1mey					1mey			٠.			Imey					lmey						Imey		
SEQ ID NO:	98 .					98					86						98					98						. 98		

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PDB annotation	DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DINA)	COMPLEX (ZINC FINGER/DINA) ZINC FINGER, PROTEIN-DNA	INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DINA) ZINC FINGER, PROTFIN-DINA	INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	FINGER/DNA) ZINC FINGER,	INTERACTION, PROTEIN	DESIGN, 2 CRYSTAL	STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC TRICED F	FINGERIDINA) ZINC FINGER, PROTEIN-DNA	INTERACTION, PROTEIN	STRUCTURE, COMPLEX	COMPLEX (ZINC	FINGER/DNA) ZINC FINGER,	INTERACTION, PROTEIN	CLER V	STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC
Compound		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN: CHAIN: C. P. G:		TAYA CHAME A D. D.	CONSENSUS ZINC FINGER	TROTEIN, CIRMIN: C, r, G,			DNA; CHAIN: A, B, D, E;	PROTEIN; CHAIN: C, F, G,			DNA: CHAIN: A B D B	CONSENSUS ZINC FINGER	FROIEIN; CRAIN: C, F, G;			DNA; CHAIN: A, B, D, B;;
SEQ FOLD score	·																			
PMF		0.36	·	0.28		700	0.00				0.70			· -	0.19					0.34
Verify score		-0.31		-0.11	·	200	900				-0.02				0.18		•			-0.25
Psi Blast		8.5e-41		1.5e-42		07 70 7	0.06-49				3.4e-48				1.7e-10					5.le-13
AA END		785		842		070	2				668				152					232
START AA		902		992		100	<u> </u>				817				123					205
CHAIN	·	ပ		၁		Ç	ر				ر ان				C)				O
PDB TD		lmey		lmey			illey				lmey				lmev					1mey
SEQ ID NO:		98		98		70	8				86				98					98

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		T			rusor.	: Oleea
PDB annotation	FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2-CRYSTAL STRUCTURE, COMPLEX CTING FINGER, DNA)	COMPLEX (ZINC FINGER, PROTEIN-DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX CANCER COMPLEX CANCER CA	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	ZINC FINGER TRANSCRIPTION FACTOR SP1; ZINC FINGER, TRANSCRIPTION ACTIVATION, SP1	COMPLEX (TRANSCRIPTION, REGULATION/DNA) TFIIIA, 5S GENE, NMR, TFIIIA, PROTEIN, DNA, TRANSCRIPTION FACTOR, 5S RNA 2 GENE, DNA BINDING PROTEIN, ZINC
Compound	CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	SPIF2; CHAIN: NULL;	TRANSCRIPTION FACTOR IIIA; CHAIN: A; SS RNA GENE; CHAIN: B, F;
SEQ FOLD score						
PMF		0.17	0.65	0.10	0.03	0.88
Verify		0.27	0.22	-0.32	-0.38	-0.07
Psi Blast		6.8e-13	6.8e-11	3.46-12	Ie-07	3.4e-21
AA END		870	668	974	873	260
START		843	871	947	846	180
CHAIN		ප	ტ .	_ව		₹
PDB UD		1mey	Ітеу	Ітеу	1sp2	£4 -
SEQ ID NO:		8	98	98	98	98

PDB annotation	FINGER, COMPLEX 3 (TRANSCRIPTION REGULATIONDNA)	COMPLEX (TRANSCRPTION REGULATION/DNA) TFIIIA; SS GENE; NMR, TFIIIA,	PROTEIN, DNA, TRANSCRIPTION FACTOR, SS RNA 2 GENE, DNA	BINDING PROTEIN, ZINC FINGER, COMPLEX 3 (TRANSCRIPTION	REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA)	REGULATION/DNA), RNA	POLYMERASE III, 2	I KANSCRIFTION INTTATION, ZINC FINGER	PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION)	POI VMERASE III 2	TRANSCRIPTION	INITIATION, ZINC FINGER	COMPLEX (TRANSCRIPTION)	REGULATION/DNA) YING-	YANG 1; TRANSCRIPTION	HINTIATION, INITIATION	FINGER PROTEIN, DNA-	PROTEIN RECOGNITION, 3 1-	COMPLEX (TRANSCRIPTION)	COMPLEX (TRANSCRIPTION)	Al
Compound		TRANSCRIPTION FACTOR IIIA; CHAIN: A; SS RNA GENE; CHAIN: E, F;				TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE;	CHAIN: B, C, E, F;				TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE;	CHAIN: B, C, E, F;				YY1: CHAIN: C: ADENO-	ASSOCIATED VIRUS P5	INITIATOR ELEMENT	DNA; CHAIN: A, B;				YY1: CHAIN: C: ADENO-	4 4 4 5 Cest man 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1
SEQ FOLD score							·																	
PMF score		0.13				0.47					0.18					0.59							0.25	
Verify		-0.32				-0.37					-0.21					-0.11							-0.51	
Psi Blast		5.1e-18				3.4e-33					le-35			-		5.1e-32							5.16-29	
END AA		785				828					£86					232				_			391	
START AA		707				707					818					135							270	
CHAIN		¥				∢			-		Ą					C	,						ن	,
PDB CD		භා				1466					146					Jubd							1.1hd	
SEQ ID NO:		98				98					98					98	1						86	3

	·	<u> </u>			
PDB annotation	REGULATION/DNA) YING- YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA- PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCREPTION REGULATION/DNA) YING- YANG 1; TRANSCREPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA- PROTEIN RECOGNITION, 3 COMPLEX (TRANSCREPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 TO COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION INITIATION ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION DNA) YING- PLYANG 1; TRANSCRIPTION PRINTIATION, INITIATIOR
Compound	ASSOCIATED VIRUS PS INTTATOR ELEMENT DNA; CHAIN: A, B;	YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS PS INTTATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;
SEQ FOLD score					·
PMF		0.00	0.45	0.83	0.18
Verify		-0.49	-0.26	-0.41	-0.35
Psi Blast		5.1e-35	3.4e-33	1.70-31	le-31
AA BA		733	763	814	668
START AA		030	658	711	796
CHAIN		O .	υ	υ _.	ပ
PDB UB		1ubd	lubd	lubd	lubd
SEQ ID NO:	·	98	98	98	98

Table 5

PDB annotation	ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA- PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING- (YANG 1; TRANSCRIPTION INTIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	TRANSCRIPTION REGULATION TRANSCRIPTION REGULATION, ADRI, ZINC FINGER, NMR	TRANSCRIPTION REGULATION TRANSCRIPTION REGULATION, ADRI, ZINC FINGER, NMR	TRANSCRIPTION REGULATION TRANSCRIPTION REGULATION, ADRI, ZINC FINGER, NMR		COMPLEX (DNA-BINDING F- PROTEIN/DNA) FIVE- FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- FI
Compound		YY1; CHAIN: C; ADENO-ASSOCIATED VRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	ADR1; CHAIN: NULL;	ADR1; CHAIN: NULL;	ADR1; CHAIN: NULL;	COMPLEX(TRANSCRIPTIO N REGULATION/DNA) TRAMTRACK PROTEIN (TWO ZINC-FINGER PEPTIDE) COMPLEXED WITH 2DRP 3 DNA 2DRP 4	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;
SEQ FOLD score			÷				
PMF score		0.13	0.80	0.22	0.10	0.07	0.89
Verify.		-0.41	-0.62	-0.10	-0.29	-0.06	-0.04
Psi Blast		6.8e-32	5.1e-13	1.2e-12	5.1e-15	1e-08	1.5e-32
END		974	289	765	901	868	260
START AA		853	237	707	.846	844	135
CHAIN		U				¥	A
PDB		Inbd .	2adr	2adr	2adr	2drp	2gli
SEQ ID		98	. 98	98	98	98	98

Table

	Т				:	4	مستعند			7			·						7			Į Ų	144 THE				31		E I	and the same	D	1			ŀ
PDB annotation	BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING	PROTEIN/DINA) FIVE-	FINGER GLL; GLL, ZINC	BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING	PROTEIN/DNA) FIVE-	FINGER GLI; GLI, ZINC	FINGER, COMPLEX (DNA-	BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING	FROI BINDER (FI. F. C. I. ZINC	FINGER, COMPLEX (DNA-	BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING	PROTEIN/DNA) FIVE-	FINGER GLI; GLI, ZINC	FINGER, COMPLEX (DNA-	BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING	PROTEIN/DINA) FIVE-	PRINCES CLI, CLI, CHINC	BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING	PROTEIN/DNA) FIVE-	FINGER GLI; GLI, ZINC	FINGER, COMPLEX (DNA-	BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING	PROTEIN/DNA) FIVE-	FINGER GLI, GLI, ZINC	BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING	PROTEIN/DNA) FIVE- FINGER GLI; GLI, ZINC	
Compound		ZINC FINGER PROTEIN	GLII; CHAIN: A; DNA;	CHAIN: C, D;		ZINC FINGER PROTEIN	GLII; CHAIN: A; DNA;	CHAIN: C, D;			ZINC FINGER PROTEIN	GLII; CHAIN: A; DNA; CHAIN: C D:			ZINC FINGER PROTEIN	GLII; CHAIN: A; DNA;	CHAIN: C, D;			ZINC FINGER PROTEIN	GLII; CHAIN: A; DNA;	CHAIN: C, D;	•	ZINC FINGER PROTEIN	GLII; CHAIN: A; DNA;	CHAIN: C, D;			ZINC FINGER PROTEIN	GLII; CHAIN: A; DNA;	CHAIN: C, D;		ZINC HINGER PROTEIN	GLII; CHAIN: A; DNA; CHAIN: C. D:	
SEQ FOLD score							*					·										,			,										**************************************
PMF		0.07		-		0.33	}				0.11	·		•	0.12					0.29				0.45					0.00				020		
Verify		-0.27	٠.			-0.57	}			-	-0.05				-0.37					-0.36				-0.44					-0.28		· · ·		930	}	
Psi Blast		1.2e-21				2.40-23					1.2e-16				1.4e-31					1.7e-32				4.8e-21					8.5e-31				2 40.21	1	
END		337				285	3		,		365				393					765				899					106				833	3	
START		180				184					207		•		270	:				630				739					772				767	30	
CHAIN		A				A	4				Ą				A					Ą				V					Ą				\ \ \	ς	
PDB		2gli				2oli	1187				2gli				2gli)	٠.			2gli				2gli	<u> </u>				2gli)			201	7 R	
SEQ ID		98				86	3				98				98					98				98					98				70	0	1

rable 5

Compound PDB annotation	FINGER, COMPLEX (DNA-BINDING PROTEINDNA)	ZINC FINGER PROTEIN GLI1; CHAIN: A; DNA; PROTEIN/DNA) FIVE- CHAIN: C, D; FINGER GLI, GLI, ZINC FINGER, COMPLEX (DNA- RINDING PROTEIN/DNA)	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; PROTEIN/DNA) FIVE- FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	HEMOLIN; CHAIN: A, B; INSECT IMMUNITY INSECT IMMUNITY, LPS-BINDING, HOMOPHILIC ADHESION	HEMOLIN; CHAIN: A, B; INSECT IMMUNITY INSECT IMMUNITY, LPS-BINDING, HOMOPHILIC ADHESION	TOTA CAMBACTE TOTAL CAMBACTE		CHAIN: NULL; IMMUNOGLOBULIN FOLD, TRANSMEMBRANE.	GLYCOPROTEIN, T-CELL, 2	MHC, LIPOPROTEIN, T-CELLF SURPACE GLYCOPROTEIN	T-CELL SURFACE T-CELL SURFACE	RCD4;	CHAIN: NULL; IMMUNOGLOBULIN FOLD,	GLYCOPROTEIN, T-CELL, 2	MHC, LIPOPROTEIN, T-CELLTI STIRFACE OF VCOPROTEIN 4.	AXONIN-1; CHAIN: A; CELL ADHESION NEURAL	 AXONIN-1; CHAIN: A; CELL ADHESION NEURAL	FIBROBLAST GROWTH GROWTH PACTOR/GROWTH! FACTOR 2; CHAIN: A, B; FACTOR RECEPTOR FGF, fl
SEQ FOLD score																		
PMR score		0.70	0.27	60:0	0.16	26.0	000				0.29					0.36	0.29	0.22
Verify score	•	0.11	-0.31	0.06	-0.07	200	† ?				0.18					-0.11	-0.18	-0.33
Psi Blast		2.4e-20	8.5e-31	3.6e-21	2.4e-11	2 40 11	7:40-11				8.4e-09					1.1e-16	2.4e-09	2.4e-11
AA A		931	976	518	962	27.6	2 1				250					518	962	250
START		818	825	23	275	150	3				30					22	284	104
CHAIN		V	A	V	¥											¥	Ą	ပ
PDB UD		2gli	2gli	1bih	1bih	1	îngr	•			lcdy					1cs6	1cs6	lcvs
SEQ ID NO:		98	98	87	87	67					87					87	28	87

Fable 5

PDB annotation	FGFR, IMMUNOGLOBULIN- LIKE, SIGNAL TRANSDUCTION, 2 DIMERIZATION, GROWTH FACTOR/GROWTH FACTOR	GROWTH FACTOR/GROWTH PACTOR RECEPTOR FGF, FGFR, IMMUNOGLOBULIN- LIKE, SIGNAL TRANSDUCTION, 2 DIMERIZATION, GROWTH FACTOR/GROWTH FACTOR RECEPTOR	CELL ADHESION NCAM; NCAM, IMMUNOGLOBULIN FOLD, GLYCOPROTEIN	CELL ADHESION NCAM; NCAM, IMMUNOGLOBULIN FOLD, GLYCOPROTEIN	CELL ADHESION NCAM; NCAM, IMMUNOGLOBULIN FOLD, GLYCOPROTEIN	CELL ADHESION NCAM; TO NCAM, IMMUNOGLOBULING FOLD, GLYCOPROTEIN	GROWTH FACTOR/GROWTH FACTOR RECEPTOR FGF2; FGFR2; IMMUNOGLOBULING (IG)LIKE DOMAINS BELONGING TO THE I-SET 2 SUBGROUP WITHIN IG-LIKE DOMAINS, B-TREFOIL FOLD.	GROWTH FACTOR/GROWTH FACTOR RECEPTOR FGF2; FGFR2; IMMUNOGLOBULIN (IG)LIKE DOMAINS BELONGING TO THE I-SET A SUBGROUP WITHIN IG-LIKEN
Compound	FIBROBL AST GROWTH FACTOR RECEPTOR 1; CHAIN: C, D;	FIBROBLAST GROWTH FACTOR 2; CHAIN: A, B; FIBROBLAST GROWTH FACTOR RECEPTOR 1; CHAIN: C, D;	NEURAL CELL ADHESION MOLECULE; CHAIN: A, B, C, D;	NEURAL CELL ADHESION MOLECULE; CHAIN: A, B, C, D;	NEURAL CELL ADHESION MOLECULE; CHAIN: A, B, C, D;	NEURAL CELL ADHESION MOLECULE; CHAIN: A, B, C, D;	FIBROBLAST GROWTH FACTOR 2; CHAIN: A, B, C, D; FIBROBLAST GROWTH FACTOR RECEPTOR 2; CHAIN: E, F, G, H;	FIBROBLAST GROWTH FACTOR 2; CHAIN: A, B, C, D; FIBROBLAST GROWTH FACTOR RECEPTOR 2; CHAIN: B, F, G, H;
SEQ FOLD score								
PMF score		0.29	0.25	0.49	0.07	0.18	0.15	0.17
Verify score		0.06	0.11	-0.03	0.05	-0.18	-0.25	-0.29
Psi Blast		2.4e-12	1.2e-10	1.2e-07	3.6e-10	8.4e-05	3.6e-11	9.6e-09
AA END		376	376	269	541	658	376	250
START AA	·	145	153	24	284	471	154	104
CHAIN		υ	¥	Ą	Ą	¥	B	ల
202 ED ED		lcvs	lepf	lepf	lepf	lepf	1ev2	lev2
SEQ ID NO:	·		87	87		87	<i>L</i> 8	87

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· PDB annotation	DOMAINS, B-TREFOIL FOLD	GROWTH FACTOR/GROWTH	FACTOR RECEPTOR FGF2;	FGFK2; IMMUNOGLOBULIN	(IG)LIKE DOMAINS	BELONGING TO THE L-SET 2	SUBGROUP WITHIN IG-LIKE	DOMAINS, B-TREFOIL FOLD	GROWTH FACTOR/GROWTH	FACTOR RECEPTOR FGF1;	FGFR1; IMMUNOGLOBULIN	(IG) LIKE DOMAINS	BELONGING TO THE I-SET 2	SUBGROUP WITHIN IG-LIKE	DOMAINS, B-TREFOIL FOLD	GROWTH FACTOR/GROWTH	FACTOR RECEPTOR FGF1;	FGFR1; IMMUNOGLOBULIN	(IG) LIKE DOMAINS	BELONGING TO THE I-SET 2	SUBGROUP WITHIN IG-LIKE	DOMAINS, B-TREFOIL FOLD	IMMUNE SYSTEM FC-	BPSILON RI-ALPHA;	IMMUNOGLOBULIN FOLD,	GLYCOPROTEIN,	RECEPTOR, IGE-BINDING 2	PROTEIN	IMMUNE SYSTEM FC- HEREN ON PLATENTY	TOWNS OF THE PARTY	CALVACODO CHODO LINE TO LOS TO THE COST OF THE COST OST OF THE COST OF THE COST OF THE COST OF THE COST OST OF THE COST OF THE COST OST OST OST OST OST OST OST OST OST	GLICOLNOIBIN,	RECEPTOR, IGE-BINDING 2 &	PROTEIN CONTRACTOR	MIMONE SYSTEM HIGH	RECEPTOR ECIEPSILON II	IGE-FC; IMMUNOGLOBULIN	FOLD, GLYCOPROTEIN,
Compound		FIBROBLAST GROWTH	FACTOR 2; CHAIN: A, B, C,	D; FIBROBLAST GROWI'H	FACTOR RECEPTOR 2;	CHAIN: E, F, G, H;			FIBROBLAST GROWTH	FACTOR 1: CHAIN: A. B.	FIBROBLAST GROWTH	FACTOR RECEPTOR 1;	CHAIN: C, D;		•	FIBROBLAST GROWTH	FACTOR 1; CHAIN: A, B;	FIBROBLAST GROWTH	FACTOR RECEPTOR 1;	CHAIN: C, D;			HIGH AFFINITY	IMMUNOGLOBULIN	EPSILON RECEPTOR	CHAIN: A;			HIGH AFFINITY	EDEN ON DECEMBE	CHAIN. A.	CHAIN: A;		A SALLAN AND THE ACT AND ASSAULT	HIGH AFFINITY	FPSH ON RECEPTOR	CHAIN: A; IG EPSILON	CHAIN C REGION; CHAIN:
SEQ FOLD score											-																											
PMF		0.24							0.05		-					0.00							90.0			,			0.31	•				200	0.23			
Verify		-0.10							-0.54	•	•					-0.08							-0.24					1	80.0					,	ې 9. آغ		:	
Psi Blast		4.8e-10							7.2e-08							8.4e-09							1.2e-08			-			2.4e-05		-			. 0000	0.00024			
END AA	·	376							250							376							250						376					,;;	541	•	· .	
START AA		153							104							143							104						154					, 8	784			
CHAIN		ß							၁					-		O							A						¥.		•	_			∢			
PDB		lev2							levt							levt							1f2q						21 P21			•		3	Itoa			
SEQ ID NO:		87							28							28							87						87						28			

						1 11111 11
PDB annotation	RECEPTOR, IGE-BINDING 2 PROTEIN, IGE ANTIBODY, IGE-FC	IMMUNE SYSTEM, MEMBRANE PROTEIN CD32; FC RECEPTOR, IMMUNOGLOULIN, LEUKOCYTE, CD32	IMMUNE SYSTEM HLA-DRI, DRA; HLA-DRI, DRBI 0101; TCR HAI.7 ALPHA CHAIN; TCR HAI.7 BETA CHAIN; PROTEIN-PROTEIN COMPLEX, IMMUNOGLOBULIN FOLD	ED,	OTEIN URAL PROTEIN, E, ON, IN FOLD, 2 ESION	IMMUNE SYSTEM
Compound	В, D;	FC RECEPTOR FC(GAMMA)RIIA; CHAIN: A;	HLA CLASS II HISTOCOMPATIBILITY ANTIGEN, DR CHAIN: A; HLA CLASS II HISTOCOMPATIBILITY ANTIGEN, DR-1 CHAIN: B; HEMAGGLUTININ HA! PEPTIDE CHAIN; CHAIN: C; T-CELL RECEPTOR ALPHA CHAIN; CHAIN: D; T-CELL RECEPTOR CHAIN; CHAIN: B;	MYELIN PO PROTEIN; CHAIN: NULL;	MYELIN PO PROTEIN; CHAIN: NULL;	SIALOADHESIN; CHAIN:
SEQ FOLD score						
PMF score		0.06	0.88	0.01	0.55	0.46
Verify score		-0.06	0.27	-0.24	0.43	90.0
· Psi Blast		0.0024	0.0048	8.4e-06	6e-05	0.00036
END AA		541	383	120	517	377
START AA		282	284	23	417	283
CHAIN		Ą	Ω		· .	A
PDB TD		Ifeg	lfyt	Ineu	Ineu	Iqfo
SEQ ID NO:		87		87		87

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PDB annotation	IMMUNOGLOBULIN SUPERFAMILY, CARBOHYDRATE BINDING	GLYCOPROTEIN CD4; IMMUNOGLOBULIN FOLD, TRANSMEMBRANE, GLYCOPROTEIN, T-CELL, 2 MHC LIPOPROTEIN, POLYMORPHISM	GLYCOPROTEIN CD4; IMMUNOGLOBULIN FOLD, TRANSMEMBRANE, GLYCOPROTEIN, T-CELL, 2 MHC LIPOPROTEIN, POLYMORPHISM	IMMUNE SYSTEM CD32; RECEPTOR, PC, CD32, IMMUNE SYSTEM	IMMUNE SYSTEM CD32; RECEPTOR, PC, CD32, IMMUNE SYSTEM	CELL ADHESION NCAM DOMAIN 1; CELL ADHESION, GLYCOPROTEIN, HEPARIN-F BINDING, GPI-ANCHOR, 2 NEURAL ADHESION MOLECULE, IMMUNOGLOBULIN FOLD, SIGNAL	CELL ADHESION NCAM DOMAIN 1; CELL ADHESION, GLYCOPROTEIN, HEPARIN-C BINDING, GPI-ANCHOR, 2 NEURAL ADHESION MOLECULE, IMMUNOGLOBULIN FOLD, F SIGNAL
Compound	A, B, C;	T-CELL SURFACE GLYCOPROTEIN CD4; CHAIN: A, B;	T-CELL SURFACE GLYCOPROTEIN CD4; CHAIN: A, B;	FC GAMMA RIB; CHAIN: A;	FC GAMMA RIB; CHAIN: A;	NEURAL CELL ADHESION MOLECULE; CHAIN: NULL;	NEURAL CELL ADHESION MOLECULE; CHAIN: NULL;
SEQ FOLD score					•		
PMF score		0.52	0.10	0.11	0.04	0.28	0.22
Verify score		0.06	-0.13	-0.11	-0.24	-0.06	0.28
Psi Blast		6e-10	6e-14	2.4e-10	0.0012	0.0024	90-99
END		486	380	250	518	119	401
START AA		160	30	104	284		284
CHAIN		∢	¥	¥	· V		
PDB		lwio	1 wio	2fcb	2fcb	2ncm	2ncm
SEQ ID NO:		87	28	87	28	87	28

Table 5

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PDB annotation	ISOMERASE FKBP; ISOMERASE, ROTAMASE	ISOMERASE FKBP;	ISOMERASE, ROTAMASE	IMMUNE SYSTEM	CALCINEURIN; FKBF12,	RAPAMYCIN, COMPLEX, RYANODINE RECEPTOR	ROTAMASE (ISOMERASE)	FKBP52 OR HSP56:	ROTAMASE (ISOMERASE),	DOMAIN I (N-TERM) OF A 59	KDA, 2 FK506-BINDING	PROTEIN, PEPTIDYL	PROLYL CIS-TRANS	ISOMERASE	ROTAMASE (ISOMERASE)	FKBP52 OR HSP56;	ROTAMASE (ISOMERASE),	DOMAIN I (N-TERM) OF A 59	KDA, 2 FK506-BINDING	PROTEIN, PEPTIDYL		ISOMERASE	. 1	CYTOKINE EBP;	_	₹		CYTOKINE 2 RECEPTOR	T COURT		ĄĽ	LICAL			יים יים יים יים יים יים יים יים יים יים
Compound	FK506 BINDING PROTEIN; CHAIN: NULL;	FK506 BINDING PROTEIN;	CHAIN: NULL;	FKBP12.6; CHAIN: A;			FKRP59-I. CHAIN: NIII I.	1.1.1.1.1.1.1.1.1.1.1.1.1.1.1.1.1.1.1.							FKBP59-I; CHAIN: NULL;	•	-							ERYTHROPOIETIN	RECEPTOR; CHAIN: A, B;				DI ACIDAMIAY I ACTIONISIE	CHAIN: A: PROLACTIN	RECEPTOR; CHAIN: B, C;				
SEQ FOLD score	95.53						93.58	2	,										-		-	-					•								
PMF score		1.00	,	1.00		•						•			1.00		•							0.45					21.0	0.10					
Verify score		0.99		0.82						•				-	0.82									-0.20					0.47	-0.4/					
Psi Blast	8.4e-44	8.4e-44		8 4 4			3.6e-41	÷							3.6e-41									2.4e-05					2 42 05	2.46-05				•	
A END	163	163		163			164	5							163									737					752	5				٠,	
START AA	95	09		57			50	3							56									. 599			-	-	777	98					
CHAIN				∢								,												. ∀					P	9				·	
PDB CI	1bkf	1bkf		legh H			Irot								Irot									lem					1666	<u> </u>					
SEQ ID NO:	06	06		S .			8	2																25						*					

Table 5

PDB annotation	DOMAINS, CYTOKINE- RECEPTOR COMPLEX		TRANSFERASE CPT; KINASE, ANTIBIOTIC RESISTANCE, PHOSPHORYLATION, 2 MONONUCLEOTIDE REMAINER FOLLOTION	PROTEIN TRANSPORT HELIX-TURN-HELIX TPR- LIKE REPEAT, PROTEIN TRANSPORT	INSECT IMMUNITY INSECT IMMUNITY, LPS-BINDING, HOMOPHILIC ADHESION	INSECT IMMUNITY INSECT IMMUNITY, LPS-BINDING, THE HOMOPHILIC ADHESION	T-CELL SURFACE GLYCOPROTEIN IMMUNOGLOBULIN FOLD, TRANSMEMBRANE, GLYCOPROTEIN, T-CELL, 2(f) MHC, LIPOPROTEIN, T-CELL, SUBBACH GLYCOPROTEIN, T-CELL,	CELL ADHESION NEURAL	CELL ADHESION NEURAL CELL ADHESION	GROWTH FACTOR/GROWTH PACTOR RECEPTOR FGF, [1] FGFR, IMMUNOGLOBULIN-[1]
Compound	DO RE	CELL ADHESION PROTEIN TENASCIN (THIRD FIBRONECTIN TYPE III REPEAT) 11EN 3	CHLORAMPHENICOL PHOSPHOTRANSFERASE; KII CHAIN: A; RE PHOSPHOTRANSFERASE; KII RE PHOSPHOTRANSFERASE; KIII RE PHOSPHOTRANSFERASE; KIIII	VESICULAR TRANSPORT FR PROTEIN SEC17; CHAIN: HE A; LIN	HEMOLIN; CHAIN: A, B; IMS IMS IMS	HEMOLIN; CHAIN: A, B; IM IM HO	T-CELL SURFACE GLYCOPROTEIN CD4; GLYCOPROTEIN CD	AXONIN-1; CHAIN: A; CE	AXONIN-1; CHAIN: A; CE	FIBROBLAST GROWTH GR FACTOR 2; CHAIN: A, B; FA FIBROBLAST GROWTH FIG
SEQ FOLD score		ОГЩК	<u> </u>	F	91.03	·		4	7	дцц
PMF score		0.05	0.07	-0.18		0.39	0.03	0.76	0.01	0.06
Verify		-0.52	-0.07	0.02	v	0.27	-0.18	-0.12	0.24	-0.00
Psi Blast		0.0012	0.0024	4.8e-11	1.2e-29	1.2e-29	0.00096	2.4e-30	3.6e-18	1.2e-21
END AA		732	513	1635	387	297	105	312	202	289
START AA		629	393	1360	18	30	22	29	9	109
CHAIN			∢	Ą	4	ď		A	A	ပ
80g 13		Iten	Iqhx	lqqe	1bih	1bih	lcdy	lcs6	1cs6	1cvs
SEQ ID NO:		94	95	96	25	26	76	16	16	

			<u> </u>						·
PDB annotation	LIKE, SIGNAL TRANSDUCTION, 2 DIMERIZATION, GROWTH FACTOR/GROWTH FACTOR RECEPTOR	GROWTH FACTOR/GROWTH FACTOR RECEPTOR FGF, FGFR, IMMUNOGLOBULIN- LIKE, SIGNAL TRANSDUCTION, 2 DIMERIZATION, GROWTH FACTOR/GROWTH FACTOR	COMPLEX, PC FRAGMENT, IGG, FC, RECEPTOR, CD16, GAMMA	COMPLEX CD16; 1GG1-FC COMPLEX, FC FRAGMENT, 1GG, FC, RECEPTOR, CD16, GAMMA	CELL ADHESION NCAM; '' NCAM, IMMUNOGLOBULIN'' FOLD, GLYCOPROTEIN	CELL ADHESION NCAM; CONCAM, INMUNOGLOBULING) FOLD, GLYCOPROTEIN	CELL ADHESION NCAM; III NCAM; III NCAM; INMUNOGLOBULIN FOLD, GLYCOPROTEIN	CELL ADHESION NCAM; 12 NCAM, IMMUNOGLOBULIN FOLD, GLYCOPROTEIN [1]	GROWTH FACTOR/GROWTH FACTOR RECEPTOR FGP2; Hg
Compound	FACTOR RECEPTOR 1; CHAIN: C, D;	FIBROBLAST GROWTH FACTOR 2; CHAIN: A, B; FIBROBLAST GROWTH FACTOR RECEPTOR 1; CHAIN: C, D;	LOW AFFINITY IMMUNOGLOBULIN GAMMA FC RECEPTOR CHAIN: C; FC FRAGMENT OF HUMAN IGG1; CHAIN: A, B;	LOW AFFINITY IMMUNOGLOBULIN GAMMA FC RECEPTOR CHAIN: C; FC FRAGMENT OF HUMAN IGG1; CHAIN: A, B;	NEURAL CELL ADHESION MOLECULE; CHAIN: A, B, C, D;	NEURAL CELL ADHESION MOLECULE; CHAIN: A, B, C, D;	NEURAL CELL ADHESION MOLECULE; CHAIN: A, B, C, D;	NEURAL CELL ADHESION MOLECULE; CHAIN: A, B, C, D;	FIBROBLAST GROWTH FACTOR 2; CHAIN: A, B, C,
SEQ FOLD score									
PMF		0.71	0.12	0.29	0.09	0.34	-0.01	0.49	0.55
Verify		-0.15	-0.00	0.39	0.12	0.57	0.10	0.20	0.06
Psi Blast		3.6e-14	0.0011	66-07	6e-22	2.4e-06	3.6e-08	3.6e-13	7.2e-22
END AA		190	147	302	285	105	312	205	289
START		30	10	113	118	15	215	30	107
CHAIN	;	ပ	А	Ą	А	Ą	Ą	¥	я
20g E0		lcvs	le4k	1e4k	lepf	lepf	lepf	lepf	lev2
SEQ ID NO:	·	26	. 20		26	<i>L</i> 6	16	26	26

				Salta State of		
PDB annotation	FOFFZ; IMMUNOGLOBULIN (IG)LIKE DOMAINS BELONGING TO THE I-SET 2 SUBGROUP WITHIN IG-LIKE DOMAINS, B-TREFOIL FOLD	GROWTH FACTOR/GROWTH FACTOR RECEPTOR FGF2; FGFR2; IMMUNOGLOBULIN (IG)LIKE DOMAINS BELONGING TO THE I-SET 2 SUBGROUP WITHIN IG-LIKE DOMAINS. B-TREFOIL FOLD	GROWTH FACTOR/GROWTH FACTOR RECEPTOR FGF2; FGFR2; IMMUNOGLOBULIN (IG)LIKE DOMAINS BELONGING TO THE I-SET 2 SUBGROUP WITHIN IG-LIKE DOMAINS, B-TREFOIL FOLD	GROWTH FACTOR/GROWTH FACTOR RECEPTOR FGF2; FGFR2; IMMUNOGLOBULIN (G)LIKE DOMAINS BELONGING TO THE I-SET 20 SUBGROUP WITHIN IG-LIKEDOMAINS, B-TREFOIL FOLD	GROWTH FACTOR/GROWTH FACTOR RECEPTOR FGF2; YEACTOR RECEPTOR FGF2; YEG)LIKE DOMAINS (IG)LIKE DOMAINS BELONGING TO THE I-SET SUBGROUP WITHIN IG-LIKED DOMAINS, B-TREFOIL FOLD	GROWTH FACTOR/GROWTH FACTOR RECEPTOR FGF1; FGFR1; IMMUNOGLOBULIN (IG) LIKE DOMAINS BELONGING TO THE I-SET A SUBGROUP WITHIN IG-LIKE
Compound	D; FIBROBLAST GROWTH FACTOR RECEPTOR 2; CHAIN: B, F, G, H;	HBROBLAST GROWTH FACTOR 2; CHAIN: A, B, C, D; FIBROBLAST GROWTH FACTOR RECEPTOR 2; CHAIN: B, F, G, H;	FIBROBLAST GROWTH FACTOR 2; CHAIN: A, B, C, D; FIBROBLAST GROWTH FACTOR RECEPTOR 2; CHAIN: B, F, G, H;	FIBROBLAST GROWTH FACTOR 2; CHAIN: A, B, C, D; FIBROBLAST GROWTH FACTOR RECEPTOR 2; CHAIN: B, F, G, H;	FIBROBLAST GROWTH FACTOR 2; CHAIN: A, B, C, D; FIBROBLAST GROWTH FACTOR RECEPTOR 2; CHAIN: B, F, G, H;	FIBROBLAST GROWTH FACTOR 1; CHAIN: A, B; FIBROBLAST GROWTH FACTOR RECEPTOR 1; CHAIN: C, D;
SEQ FOLD score						
PMF score		0.63	0.16	0.46	60.0	0.12
Verify score		0.01	-0.05	0.12	-0.00	-0.06
Psi Blast	,	900000	3.6e-16	1.26-24	3.6e-18	2.4e-23
END		86	190	297	204	289
START AA		1.5	29	107	29	107
CHAIN		ш	E	5	5 .	ပ
PDB ID		lev2	lev2	lev2	lev2	levt
SEQ ID NO:	·	<i>L</i> 6	97	26	16	97

Table 5

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PDB annotation	DOMAINS, B-TREFOIL FOLD	IMMUNE SYSTEM FC. EPSILON RI-ALPHA; IMMUNOGLOBULIN FOLD,	GLYCOPROTEIN, RECEPTOR, IGE-BINDING 2	PROTEIN	DAMUNE SYSTEM FC. EPSILON RI-ALPHA;	IMMUNOGLOBULIN FOLD, GLYCOPROTEIN.	RECEPTOR, IGE-BINDING 2 PROTEIN	IMMUNE SYSTEM FC.	EPSILON KI-ALPHA; IMMUNOGLOBULIN FOLD,	GLYCOPROTEIN,	RECEPTOR, IGE-BINDING A	IMMUNE SYSTEM HIGH	AFFINITY IGE-FC	RECEPTOR, FC(EPSILON) IGE-FC: IMMUNOGLOBULIN	FOLD, GLYCOPROTEIN,	2	PROTEIN, IGE ANTIBODY, 'I	IMMUNE SYSTEM HIGH	AFFINITY IGE-FC	IGE-FC; IMMUNOGLOBULIN	FOLD, GLYCOPROTEIN,	RECEPTOR, IGE-BINDING 2,	IGE-FC	IMMUNE SYSTEM, ————————————————————————————————————	FC RECEPTOR, A TURNING TO THE TRANSPORT OF THE TRANSPORT
Compound	,	HIGH AFFINITY IMMUNOGLOBULIN EPSILON RECEPTOR	CHAIN: A;		HIGH AFFINITY IMMUNOGLOBULIN	EPSILON RECEPTOR CHAIN: A:		HIGH AFFINITY	IMMUNOGLOBULIN EPSILON RECEPTOR	CHAIN: A;		HIGH AFFINITY	IMMUNOGLOBULIN	EPSILON RECEPTOR CHAIN: A: 1G EPSILON	CHAIN C REGION; CHAIN:	B, D;		HIGH AFFINITY	IMMUNOGLOBULIN FPSIT ON PHCHPTOR	CHAIN: A: IG EPSILON	CHAIN CREGION; CHAIN:	в, D;		FC RECEPTOR FC(GAMMA)RIIA; CHAIN:	A;
SEQ FOLD score				.																					
PMF score		0.71	-		0.36	,		0:30				0.72			•			0.88				•		0.22	
Verify score		0.31			0.36			0.30				0.35						0.25						0.34	
Psi Blast		1.2e-21			1.2e-08			6e-23	•			1.2e-23						4.8e-24						2.4e-25	
END		297	,		109		•	204				297						204						292	
START AA		115			91			30				108			,. <u>-</u>			16						115	
CHAIN		ď			¥			Ą				Ą						A						A	
PDB U		1229			1f2q			1f2q				1f6a						1f6a						1fcg	
SEQ ID NO:		76			26			. 46				97						26						76	

Table 5

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PDB annotation	LEUKOCYTE, CD32	IMMUNE SYSTEM, MEMBRANE PROTEIN CD32; FC RECEPTOR, IMMUNOGLOULIN, LEUKOCYTE, CD32	IMMUNE SYSTEM, MEMBRANE PROTEIN CD32; FC RECEPTOR, IMMUNOGLOULIN, LEUKOCYTE, CD32			Σ. q	OR TORY L	GLYCOPROTEIN CD4; IMMUNOGLOBULIN FOLD, F TRANSMEMBRANB, GLYCOPROTEIN, T-CELL, 2
Compound		FC RECEPTOR FC(GAMMA)RIIA; CHAIN: A;	FC RECEPTOR FC(GAMMA)RIIA; CHAIN: A;	TLYMPHOCYTE ADHESION GLYCOPROTEIN CD2 (HUMAN) 1HNF 3	T LYMPHOCYTE ADHESION GLYCOPROTEIN CD2 (HUMAN) 1HNF 3	PS8-CI 42 KIR; CHAIN: NULL;	PS8-CL42 KIR; CHAIN: NULL;	T-CELL SURFACE GLYCOPROTEIN CD4; CHAIN: A, B;
SEQ FOLD score								73.40
PMF score		0.40	69.0	0.12	0.10	0.13	0.13	
Verify		0.06	0.43	0.17	-0.03	0.33	0.20	
Psi Blast		1.16-23	8.4e-09	6e-05	3.6e-12	4.8e-15	6e-12	9.6e-14
AA END		203	297	&	187	276	201	376
START		16	208	25	30	120	20	39
CHAIN		¥	Y					Y
PDB UD		1fcg	lfcg	lbaf	lbuf	lnkr	lnkr	Iwio
SEQ ID NO:		97	97	97			<i>16</i>	97

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PDB annotation	MHC LIPOPROTEIN, POLYMORPHISM	IMMUNE SYSTEM CD32; RECEPTOR, PC, CD32, IMMUNE SYSTEM	IMMUNE SYSTEM CD32; RECEPTOR, FC, CD32, IMMUNE SYSTEM		INSECT IMMUNITY INSECT IMMUNITY, LPS-BINDING, HOMOPHILIC ADHESION	INSECT IMMUNITY INSECT IMMUNITY, LPS-BINDING, HOMOPHILIC ADHESION	T-CELL SURFACE	GLYCOPROTEIN IMMUNOGLOBULIN FOLD,	TRANSMEMBRANE,	GLYCOPROTEIN, T-CELL, 2	SURFACE GLYCOPROTEIN	CELL ADHESION NEURAL	CELL AUTESION	GROWTH FACTOR/GROWTH FACTOR RECEPTOR FGF,	FGFR, IMMUNOGLOBULIN-		TRANSDUCTION, 2	DIMERIZATION, GROWTH	RECEPTOR	COMPLEX CD16; IGG1-FC	COMPLEX, FC FKAGMEN 1.	· cath		COMPLEX CD16; IGG1-FC	
Compound		FC GAMMA RIB; CHAIN: A;	FC GAMMA RIB; CHAIN: A;		HEMOLIN; CHAIN: A, B;	HEMOLIN; CHAIN: A, B;	T-CELL SURFACE	GLYCOPROTEIN CD4; CHAIN: NULL;				AXONIN-1; CHAIN: A;	TOWNER OF THE LANGE CONTRACTOR	FIBROBLAST GROWTH FACTOR 2; CHAIN: A, B;	FIBROBLAST GROWTH	FACTOR RECEPTOR 1;	CHAIN: C, D;			LOW AFFINITY	CANNA HO PROPERTY	CHAIN: C. HO HE AGMENT	OF HUMAN IGG1; CHAIN: A. B:	LOW AFFINITY	
SEQ FOLD score		·			86.68						,											·			
PMF		0.64	0.49			0.39	0.03					0.76	100	0.00						0.29				0.46	
Verify		0.23	0.23			0.27	-0.18					-0.12		90.00						0.39				-0.31	
Pst Blast		1.2e-28	2.4e-22		1.2e-29	1.2e-29	96000.0					2.4e-30		1.2e-21 						6e-07				0.0036	
A END		297	204		380	290	86			÷		305	000	282						295				54	
START AA		114	16		11	23	15					22	98,	102						106				24	
CHAIN	·	¥	А		¥	A						Ą		ပ						Ą	٠.			A	
PDB U		2fcb	2fcb		1bih	1bih	1cdy					1cs6		Icvs						1e4k				1e4k	
SEQ ID NO:		26	26		86	86	86					86	8	86						86				98	

Table

PDB annotation	COMPLEX, FC FRAGMENT, IGG, FC, RECEPTOR, CD16, GAMMA	CELL ADHESION NCAM; NCAM, IMMUNOGLOBULIN FOLD, GLYCOPROTEIN	CELL ADHESION NCAM; NCAM, IMMUNOGLOBULIN FOLD, GLYCOPROTEIN	CELL ADHESION NCAM; NCAM, IMMUNOGLOBULÍN FOLD, GLYCOPROTEIN	GROWTH FACTOR/GROWTH FACTOR RECEPTOR FGF2; FGFR2; IMMUNOGLOBULIN (IG)LIKE DOMAINS BELONGING TO THE I-SET 2 SUBGROUP WITHIN IG-LIKE DOMAINS, B-TREFOIL FOLD	GROWTH PACTOR/GROWTH PACTOR RECEPTOR PGF2; FGFR2; IMMUNOGLOBULING (G)LIKE DOMAINS BELONGING TO THE 1-SET \$\vec{\vec{\vec{\vec{\vec{\vec{\vec{	GROWTH FACTOR/GROWTE FACTOR RECEPTOR FGF2; [I] FGFR2; IMMUNOGLOBULIN (IG)LIKE DOMAINS BELONGING TO THE I-SET [I] SUBGROUP WITHIN IG-LIKE; DOMAINS, B-TREFOIL FOLITE	GROWTH PACTOR/GROWTH, PACTOR RECEPTOR FGF2; [1] FGFR2; IMMUNOGLOBULIN [1] (IG)LIKE DOMAINS [1]
Compound	IMMUNOGLOBULIN GAMMA FC RECEPTOR CHAIN: C; FC FRAGMENT OF HUMAN IGG1; CHAIN: A, B;	NEURAL CELL ADHESION MOLECULE; CHAIN: A, B, C, D;	NEURAL CELL ADHESION MOLECULE; CHAIN: A, B, C, D;	NEURAL CELL ADHESION MOLECULE; CHAIN: A, B, C, D;	FIBROBLAST GROWTH FACTOR 2; CHAIN: A, B, C, D; FIBROBLAST GROWTH FACTOR RECEPTOR 2; CHAIN: E, F, G, H;	FIBROBLAST GROWTH FACTOR 2; CHAIN: A, B, C, D; FIBROBLAST GROWTH FACTOR RECEPTOR 2; CHAIN: E, F, G, H;	FIBROBLAST GROWTH FACTOR 2; CHAIN: A, B, C, D; FIBROBLAST GROWTH FACTOR RECEPTOR 2; CHAIN: B, F, G, H;	FIBROBLAST GROWTH FACTOR 2; CHAIN: A, B, C, D; FIBROBLAST GROWTH FACTOR RECEPTOR 2;
SEQ FOLD score				·				
PMF score		60:00	0.46	-0.01	0.55	0.48	0.16	0.46
Verify score		0.12	0.35	0.10	0.06	90.0-	-0.05	0.12
Psi Blast		6e-22	4.8e-06	3.6e-08	7.2e-22	0.0012	3.6e-16	1.2e-24
END		278	86	305	282	91	183	290
START		111	17	208	001	18	22	100
CHAIN		A	A	A	ш	a	ы	ප
PDB TO		lepf	lepf	lepf	lev2	lev2	lev2	lev2
SEQ ID	·	86	86	86	86 .	86	86	86

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Compound PDB annotation	F, G, H; SUBGROUP WITHIN IG-LIKE DOMAINS, B-TREFOIL FOLD	-	FACTOR 2; CHAIN: A, B, C, FACTOR RECEPTOR FORZ; D; FIBROBLAST GROWTH FGFR2; IMMUNOGLOBULIN	OR 2;		SUBGROUP WITHIN IG-LIKE DOMAINS, B-TREFOIL FOLD	FIBROBLAST GROWTH GROWTH FACTOR/GROWTH) -	FIOK I:	O; SELCINGING TO THE 1-SEL 2	DOMAINS B-TREFOIL FOLD				GLYCOPROTEIN,	RECEPTOR, IGE-BINDING 2	NITY IMMUNE SYSTEM FC.			RECEPTOR IGE-RINDING 2 1/2	PROTEIN				CHAIN: A; IG EPSILON IGB-FC; IMMUNOGLOBULING	_		IGB-FC	IMMUNE SYSTEM HIGH AFFINITY IGE-RC	
ပိ	CHAIN: B, F, G, H;	FIBROBLA	D; FIBROB	FACTOR R	CHAIN: E, F, G, H;		FIBROBLA.	PACTOR 1;	FIBROBLA	HACIOKK	CHAIN: C. D.		HIGH AFFINITY	IMMUNOGLOBULIN	EPSILON RECEPTOR	CHAIN: A;	•	HIGH AFFINITY	DONOMINI TIME	CHAIN: 4:			HIGH AFFINITY	IMMUNOGLOBULIN	EPSILON RECEPTOR	CHAIN: A:	Target Ch	֓֞֜֞֜֜֜֝֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֓֡֓֓֓֓֓֓֓֡֓֓֓֡		HIGH AFFINITY IMMUNOGLOBULIN	
SEQ FOLD score									-						•																L
PMF		60.0					0.12						0.71					0.30					0.72						_	0.13	
Verify score		-0.00					-0.06						0.31					0:30					0.35							0.33	
Psi Blast	·	3.6e-18					2.4e-23	-	-				1.2e-21	_				6e-23					1.2e-23							7.2e-24	
END		197					282						290					197	-				290							197	
START AA		22					100						108					23					101						•	14	
CHAIN		д					ပ						V					¥			- -		Ą							¥	
PDB EDB		1ev2					levt						1220	•				b731					1f6a							1f6a	
SEQ ID NO:		86					86						86					86					86							86	

Table

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PDB annotation	RECEPTOR, FC(EPSILON) IGE-FC; IMMUNOGLOBULIN FOLD, GLYCOPROTBIN, RECEPTOR, IGE-BINDING 2 PROTEIN, IGE ANTIBODY, IGE-FC	IMMUNE SYSTEM, MEMBRANE PROTBIN CD32; FC RECEPTOR, IMMUNOGLOULIN, LEUKOCYTE, CD32	IMMUNE SYSTEM, MEMBRANE PROTEIN CD32; FC RECEPTOR, IMMUNOGLOULIN, LEUKOCYTE, CD32	IMMUNE SYSTEM, MEMBRANE PROTEIN CD32; FC RECEPTOR, IMMUNOGLOULIN, LEUKOCYTE, CD32	IMMUNE SYSTEM, MEMBRANE PROTEIN CD32; FC RECEPTOR, IMMUNOGLOULIN, LEUKOCYTE, CD32		Hindi Unase et 1	INHIBITORY RECEPTOR KILLER CELL INHIBITORY RECEPTOR, INHIBITORY RECEPTOR, NATURAL KILLER CELLS,
Compound	EPSILON RECEPTOR CHAIN: A; IG EPSILON CHAIN C'REGION; CHAIN: B, D;	FC RECEPTOR FC(GAMMA)RIIA; CHAIN: A;	FC RECEPTOR FC(GAMMA)RIIA; CHAIN: A;	FC RECEPTOR FC(GAMMA)RIIA; CHAIN: A;	FC RECEPTOR FC(GAMMA)RIIA; CHAIN: A;	TLYMPHOCYTE ADHESION GLYCOPROTEIN CD2 (HUMAN) IHNF 3	T LYMPHOCYTE ADHESION GLYCOPROTEIN CD2 (HUMAN) 1HNF 3	PS8-CI 42 KIR; CHAIN: NULL;
SEQ FOLD score								
PMF score		0.22	0.10	0.10	0.69	0.12	0.10	0.13
Verify		0.34	0.49	0.05	0.43	0.17	-0.03	0.33
Psi Blast		2.4e-25	1.2e-06	2.4e-23	8.4e-09	6e-05	3.6e-12	4.8e-15
END AA		285	66	196	290	83	180	269
START AA		108	12	17	201	18	23	113
CHAIN		4	∢	∢	∢			
EQE EA		1fcg	1fcg	lfcg	Ifcg	lhaf	lhaf	1nkr
SEQ ID NO:		86	86	86	86	86	86	86

PDB annotation	IMMUNOLOGICÁL 2 RECEPTORS, IMMUNOGLOBULIN FOLD	GLYCOPROTEIN CD4; IMMUNOGLOBULIN FOLD, TRANSMEMBRANE, GLYCOPROTEIN, T-CELL, 2 MHC LIPOPROTEIN, POLYMORPHISM	IMMUNE SYSTEM CD32; RECEPTOR, PC, CD32, IMMUNE SYSTEM	IMMUNE SYSTEM CD32; RECEPTOR, FC, CD32, IMMUNE SYSTEM	BLOOD CLOTTING PROTEIN INHIBITOR COMPLEX, COAGULATION COPACTOR, PROTEASE	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING (THOGER, DNA-BINDING (THOGER)	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC (I) FINGER, DNA-BINDING (I) PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING FINGER, PROTEIN
Compound		T-CELL SURFACE GLYCOPROTEIN CD4; CHAIN: A, B;	FC GAMMA RIIB; CHAIN: A;	FC GAMMA RIB; CHAIN: A;	COAGULATION FACTOR XA; CHAÎN: A; COAGULATION FACTOR XA; CHAÎN: L;	OGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B,	QGSR ZINC FINGER PEPTIDE, CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDB; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;
SEQ FOLD score	·	72.45						
PMF score			0.64	0.45	0.00	0.74	1.00	1.00
Verify score	•		0.23	0.34	0.23	-0.09	0.27	0.64
Psi Blast		9.6e-14	1.2e-28	4.8e-22	0.00072	4.8e-27	1.1e-37	7.26-42
AA BAD		369	290	197	1104	260	288	316
START AA		32	107	17	1068	182	209	235
CHAIN		V	Ą	Ą	ı	A	ď	V
PDB ED		1wio	2fcb	2fcb	1fjs	lalh	laih	lalh
SEQ ID NO:		86	86	86	537	102	102	102

Table 5

				T The	page maps		
PDB annotation	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX
Compound	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUFLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	OGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX
SEQ FOLD score		·					
PMF	1.00	1.00	1.00	1.00	1.00	1.00	1.00
Verify score	0.38	0.10	0.34	0.02	0.46	-0.11	0.10
Psi Blast	6e-45	3.6e-46	7.2e-46	1.2e-44	2.4e-45	3.6e-46	2.4e-46
END AA	344	372	400	428	456	484	511
START AA	263	291	319	347	375	403	431
CHAIN	Ą	Ą	¥	¥	¥	A	A
PDB TD	laih	laih	lalh	lalh	lalh	laih	lalh
SEQ ID NO:	102	1,02 1,02	102	102	102	102	102

Table

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PDB annotation	FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN		
Compound	OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	TRANSCRIPTION REGULATION YEAST TRANSCRIPTION FACTOR ADRI (RESIDUES 102 - 130) 1ARD 3 (AMINO TERMINAL ZINC FINGER DOMAIN) (NMR, 10 STRUCTURES) 1ARD 4 (ADRIB) 1ARD 5	TRANSCRIPTION REGULATION YEAST TRANSCRIPTION FACTOR ADRI (RESIDUES 102 - 130) 1ARD 3 (AMINO TERMINAL ZINC FINGER DOMAIN) (NMR, 10 STRUCTURES) 1ARD 4 (ADRIB) 1ARD 5
SEQ FOLD						
PMF		1.00	0.39	0.71	0.94	0.77
Verify		-0.17	0.11	0.33	0.10	0.83
Psi		1.2e-31	3.6e-27	1.2e-21	2.46-10	8.46-10
END		532		649	235	514
START		459	564	592	209	487
CHAIN		«	∢ .	∢		·
PDB		laih	laih	laih	lard	lard
SEQ ID		102	102	102	102	102

			·	PET/USUE/DI	the limit for
PDB annotation				(E. 15mile (E. 1s. Januar, anda, species for stands, court, court	ZINC FINGER TRANSCRIPTION FACTOR SP1; ZINC FINGER,
Compound	DNA-BINDING PROTEIN HUMAN ENHANCER- BINDING PROTEIN MBP-1 MUTANT WITH CYS 11 1BBO 3 REPLACED BY ABU (C11ABU) (NMR, 60 STRUCTURES) 1BBO 4	DNA-BINDING PROTEIN HUMAN ENHANCER- BINDING PROTEIN MBP-1 MUTANT WITH CYS 11 IBBO 3 REPLACED BY ABU (C11ABU) (NMR, 60 STRUCTURES) 1BBO 4	DNA-BINDING PROTEIN HUMAN ENHANCER. BINDING PROTEIN MBP-1 MUTANT WITH CYS 11 1BBO 3 REPLACED BY ABU (C11ABU) (NMR, 60 STRUCTURES) 1BBO 4	TRANSCRIPTION REGULATION YEAST TRANSCRIPTION FACTOR ADRI (RESIDUES 130 - 159) IPAA 3 (PAPA - CARBOXY TERMINAL ZINC FINGER DOMAIN) MUTANT WITH IPAA 4 PRO 131 REPLACED BY ALA, PRO 133 REPLACED BY ALA, CYS 140 IPAA 5 REPLACED BY ALA (P131A,P133A,C140A) (NMR, 10 STRUCTURES)	SPIF3; CHAIN: NULL;
SEQ FOLD score					•
PMF	0.13	-0.01	0.12	66:0	0.18
Verify	-0.32	20.0	-0.36	0.38	0.15
Psi Blast	6e-27	2.4e-19	2.4e-19	4.8e-07	6e-09
END AA	516	970	648	513	514
START AA	461	266	594	487	487
CHAIN ID					·
208 30 CE	1bbo	1bbo	1 bbo	lpaa	lsp1
SEQ ID NO:	102	102	102	102	102

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PDB annotation	TRANSCRIPTION ACTIVATION, SP1	ZINC FINGER	TRANSCRIPTION FACTOR SP1; ZINC FINGER,	TRANSCRIPTION	ACTIVATION, SPI	ZINC FINGER TRANSCRIPTION FACTOR	SP1; ZINC FINGER,	TRANSCRIPTION COLUMN	ACTIVATION, SFI	TRANSCRIPTION FACTOR	SP1; ZINC FINGER,	TRANSCRIPTION	ACTIVATION, SP1	COMPLEX (TRANSCRIPTION	REGULATION/DNA) TFIIIA;	SS GENE; NMK, I'FILIA,	TROIEIN, DINA,	SC DNA 2 CENT DNA	BINDING PROTEIN ZINC	ENICED CONDITED 4	THATTEN, COMPLETE S	(IRAINSCRIPTION FREGULATION/DNA)	COMPLEX (TRANSCRIPTION)	REGULATION/DNA) TFIIIA; F	SS GENE; NMR, TFIIIA,	PROTEIN, DNA,	IKANSCRIPTION FACTOR,	SS KNA 2 GENE, DNA PINDING PROTEIN TING	TANGED CONDITION 2	THINGER, COMPLEAS	REGULATION/DNA)	COMPLEX (TRANSCRIPTION FREGITATION THINS: IT	H
Compound		SP1F2; CHAIN: NULL;				SPIF2; CHAIN: NULL;			CDITO. CITATAL MITH	OF 11'2, CATAIN, INOLLS,				TRANSCRIPTION FACTOR	IIIA; CHAIN: A; 58 RNA	GENE; CHAIN: B, F;							TRANSCRIPTION FACTOR	IIIA; CHAIN: A; 5S RNA	GENE; CHAIN: E, F;							TRANSCRIPTION FACTOR	
SEQ FOLD score																																	
PMIF score		0.59			300	0.99			0.53					0.11									0.51									0.99	
Verify		0.49			200	0.37		:	750	}				-0.05							٠	-	20.0	,			-	,				0.51	
.Psi Blast		3.6e-11			,	1.26-12			3 60 00					66-16								,	£6-27					•	•			4.8e-33	
END		319			7	459			514	;				261									290					•	٠			346	
START AA		291			107	431			487	· .	,			182								٠	500									263	
CHAIN ID											-			∢									Ą									∢	
PDB ID		1sp2			5	7ds1			CmJ	L I			1	<u> </u>									143									163	
SEQ ID NO:		102			5	701			102	}				102									102									102	

Table 5

					
PDB annotation	5S GENE; NMR, TFIIIA, PROTEIN, DNA, TRANSCRIPTION FACTOR, 5S RNA 2 GENE, DNA BINDING PROTEIN, ZINC FINGER, COMPLEX 3 (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) TFIIIA; 5S GENE; NMR, TFIIIA, PROTEIN, DNA, TRANSCRIPTION FACTOR, 5S RNA 2 GENE, DNA BINDING PROTEIN, ZINC FINGER, COMPLEX 3 (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) TFIIIA; 5S GENE; NMR, TFIIIA, PROTEIN, DNA, TFIIIA, TRANSCRIPTION PACTOR, 5S RNA 2 GENE, DNA BINDING PROTEIN, ZINC FINGER, COMPLEX 3 (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION DNA, THIIA, 5S GENB; NMR, THIIA, PROTEIN, DNA, THIIA, TRANSCRIPTION FACTOR, 5S RNA 2 GENE, DNA BINDING PROTEIN, ZINC FINGER, COMPLEX 3 (TRANSCRIPTION REGULATIONDNA)	COMPLEX (DNA-BINDING
Compound	GENB; CHAIN: B, F,	TRANSCRIPTION FACTOR IIIA; CHAIN: A; 5S RNA GENE; CHAIN: B, F;	TRANSCRIPTION FACTOR IIIA; CHAIN: A; 5S RNA GENE; CHAIN: B, F;	TRANSCRIPTION FACTOR IIIA; CHAIN: A; 5S RNA GENE; CHAIN: B, F;	ZINC FINGER PROTEIN
SEQ FOLD score					
PMF		0.41	-0.03	0.34	0.95
Verify		0.05	0.20	0.24	0.03
Psi Blast		6e-21	2.4c-15	9.6e-15	1.2e-54
END AA		541	638	649	345
START AA		459	564	592	182
CHAIN		A	· ·	V	V
PDB ID		1473	1473	11 C	2gli
SEQ ID	·	102	201	102	102

Table

					· · · · · · · · · · · · · · · · · · ·		or that would be that the	
PDB annotation	PROTEIN/DNA) FIVE- FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE- FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE- FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE- FINGER GLI, GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE- FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE. FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA-BINDING PROTEIN/DNA)	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC
Compound	GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLTI; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLI1: CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX
SEQ FOLD score				96.84				
PMF		1.00	1.00		1.00	09:0	0.25	0.03
Verify score		0.48	0.19		00:0-	0.14	-0.10	0.03
Psi Blast		2.46-70	2.4e-74	2.46-74	1.2e-73	6e-51	3.46-24	8.5e-25
END AA		373	457	457	513	618	195	222
START AA		234	290	318	346	430	123	144
CHAIN		¥	¥	Ą	Ą	A	V	Y
EGE CE		2gli	2gli	2gli	2gli	2gli	lalh	lalh
SEQ ID NO:		102	102	102	102	102	103	103

Table 5

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PDB annotation	FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, F PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX	COMPLEX (ZINC FINGER/DNA) ZINC FINGER,
Compound	OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B,	DNA; CHAIN: A, B, D, B; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E, CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, B; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, B; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, B; CONSENSUS ZINC FINGER
SEQ FOLD score		·					
PMF score		0.47	1.00	0.98	0.89	1.00	1.00
Verify score		0.03	0.13	0.16	-0.46	0.40	0.31
Psi Blast		1.2e-42	8.5e-41	3.4e-43	1.4e-43	1.7e-49	1.2e-50
END AA			195	222	293	321	349
START AA		325	120	143	198	240	268
CHAIN		∢	U	ပ	v	U	၁
PDB ID		lalh	Imey	Imey	1mey	1теу	lmey
SEQ ID NO:		103	103	103		103	103

PDB annotation	PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGHR/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX-(ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX
Compound	PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E, CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;
SEQ FOLD score		99.05			·	
PMF score			1.00	1.00	1.00	1.00
Verify score		·	0.42	0.36	0.47	0.50
Psi Blast		1.7e-51	1.7e-51	5.1e-51	16-50	6.8e-51
END AA		350	377	405	433	461
START AA		268	296	324	352	380
CHAIN		ပ	ပ	ပ	U	U
PDB ID		Imey	Ітеу	Imey	lmey	1mey
SEQ ID NO:		103	103	103	103	103

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PDB annotation	(ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER.	PROTEIN-DNA	INTERACTION, PROTEIN	DESIGN, 2 CRYSTAL	STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC	FINGER/DNA) ZINC FINGER,	PROTEIN-DNA	INTERACTION, PROTEIN	DESIGN, 2 CRYSTAL	STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC	FINGER/DNA) ZINC FINGER,	PROTEIN-DNA	INTERACTION, PROTEIN	DESIGN, 2 CRYSTAL	STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC	FINGER/DNA) ZINC FINGER,	PROTEIN-DNA	INTERACTION, PROTEIN	DESIGN, 2 CRYSTAL	STRUCTURE, COMPLEX	(ZINC FINGER/UNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA) THIIIA;	5S GENE; NMR, TFIIIA,	PROTEIN, DNA,	TRANSCRIPTION FACTOR,	5S RNA 2 GENE, DNA	EINCHO COMPLEY 3	(TRANSCRIPTION	REGULATION/DNA)
Compound		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;					DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;					DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;					DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;			- ·	The same of the same of the same	TRANSCRIPTION FACTOR	IIIA; CHAIN: A; 58 RNA	GENE; CHAIN: E, F;						
SEQ FOLD score																													,	٠.							
PMF score		1.00				•		1.00	- *				-	-	0.12	-						1.00			•			,	0.0								
Verify		0.80						0.29	•						-0.02							0.94		,		-		200	-0.23								
Psi Blast		1e-50						5.1e-43	-						3.4e-11							1.7e-12						,	1.2e-14						,		
A ES		489						2 04							168							293						125	55								
START AA		408					,	436							141							766						55	57	_	_	-					
CHAIN		ວ						ပ							ڻ ن							Ō							<								
PDB ID		Imey						lmey							1mey							lmey						27.	<u> </u>								
SEQ ID NO:		103						103							103				•			103						50.	103			-					

Table 5

PDB annotation	COMPLEX (TRANSCRIPTION REGULATION/DNA) TFIIIA; 5S GENE; NMR, TFIIIA, PROTEIN, DNA, TRANSCRIPTION PACTOR, 5S RNA 2 GENE, DNA BINDING PROTEIN, ZINC FINGER, COMPLEX 3 (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASB III, 2 TRANSCRIPTION INTIATION, ZINC FINGER	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTIONS REGULATIONDNA) COMPLEX (TRANSCRIPTION REGULATIONDNA), RNA FL POLYMERASE III, 2
Compound	TRANSCRIPTION FACTOR IIIA; CHAIN: A; SS RNA GENE; CHAIN: B, F;	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, B, F;	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;
SEQ FOLD score			100.94		·
PMF score	90'0	0.86		0.99	1.00
Verify	0.03	-0.20		60.0	0.26
Psi Blast	6.86-17	8.5e-35	5.1e-38	5.1e-38	1.7e-38
END AA	218	358	403	386	503
START AA	44.	199	239	241	353
CHAIN	∢	V	V	⋖	Ą
PDB ED	163	1116	1466	1tf6	14f6
SEQ ID NO:	103	103	103	103	103

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PDB annotation	TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR	ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA- PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION	INITIATION, INITIATOR ELEMENT, YYI, ZINC 2 FINGER PROTEIN, DNA-	PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-	YANG 1; TRANSCRIPTION INITIATION, INITIATOR	FILEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-	COMPLEX (TRANSCRIPTION)	COMPLEX (TRANSCRIPTION	YANG 1; TRANSCRIPTION	ELEMENT, YY1, ZINC2	FINGER PROTEIN, DNA- PROTEIN RECOGNITION, 3	COMPLEX (TRANSCRIPTION)	
Compound		YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;		YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT	DNA; CHAIN: A, B;		YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS P5	INITIATOR ELEMENT DNA; CHAÎN: A, B;			YY1; CHAIN: C; ADENO-	ASSOCIATED VIKUS F3 INITIATOR ELEMENT DAMA: CHAIN: A R.	DNA, CIMIN: A, B;	•		
SEQ FOLD score											84.44					
PMF		0.03		1.00			1.00									
Verify score		0.08		0.02			0.30								,	
Psi Blast		1e-29		5.1e-32			3.4e-35				3.4e-35					
END AA		223		321		-	349				378					
START AA		123		218			248				270	····				
CHAIN	·	ບ		၁			S				ပ		٠,		-	
PDB UD		lubd		lubd			lubd				lubd				•	
SEQ ID NO:		103		103			103				103					

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					- 1 - 1 - 1 - 1 - 1 - 1 - 1 - 1 - 1 - 1
PDB annotation	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INTIATION INTIATION ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING- YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 HINGER PROTEIN, DNA- PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 FINGER (TRANSCRIPTION) REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT;-YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, SEGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION
Compound	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR HIEMENT DNA; CHAIN: A; B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT
SEQ FOLD score					
PMF score	1.00	1.00	1.00	1.00	1.00
Verify score	0.21	0.26	0.40	0.16	0.28
Psi Blast	8.5e-35	3.4e-35	1.5e-34	3.4e-35	1.5e-29
END	377	433	461	489	504
START AA	276		360	388	416
CHAIN	ပ	O .	ပ	၁	၁
PDB ID	Iubd	lubd	Iubd	lubd	Iubd
SEQ ID NO:	103	103		103	103

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PDB annotation	INITIATION, INITIATOR ELEMENT, YYI, ZINC 2	FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3	COMPLEX (TRANSCRIPTION REGULATION/DNA)	TRANSCRIPTION	TRANSCRIPTION	REGULATION, ADRI, ZINC FINGER, NMR	TRANSCRIPTION	TRANSCRIPTION	REGULATION, ADRI, ZINC FINGER NAR	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE.	FINGER GLI; GLI, ZINC	FINGER, COMPLEX (DNA-	BINDING FROI EIN/DINA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-	FINGER GLI, GLI, ZINC	FINGER, COMPLEX (DNA-	COMPLEX (DNA-BINDING	PROTEIN/DNA) FIVE-	FINGER GLI; GLI, ZINC	FINGER, COMPLEX (DNA-	COMPLEX (DNA-BINDING F	PROTEIN/DNA) FIVE-	FINGER GLI, GLI, ZINC	BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING	PROTEIN/DNA) FIVE- FINGER GLI; GLI, ZINC
Compound	DNA; CHAIN: A, B;			ADR1; CHAIN: NULL;			ADRI; CHAIN: NULL;			ZINC FINGER PROTEIN	CHAIN: C, D;			ZINC FINGER PROTEIN GLII; CHAÎN: A; DNA;	CHAIN: C, D;		ZINC FINGER PROTEIN	GLII; CHAIN: A; DNA;	CHAIN: C, D;		ZINC FINGER PROTEIN	GLI1; CHAIN: A; DNA;	CHAIN: C, D;		ZINC FINGER PROTEIN	GLII; CHAIN: A; DNA; CHAIN: C, D;
SEQ FOLD score																					92.19					
PMF				00.00			0.17			0.23				0.90			1.00					•			1.00	
Verify score				-0.14			0.02			-0.40				0.17			0.51								0.29	
Psi Blast				3.4e-13			1.7e-13			3.6e-52				1.7e-33			1.2e-66			•	1.2e-70				1.2e-70	
END	·			170			197			351				348			379				407				433	
START AA	·			120			144			143				205			244				268				. 892	
CHAIN										Ą				∢			A				A				A	
PDB UU				2adr			2adr			2gli				2gli			2gli				2gli				2gli	
SEQ ID NO:				103			103			103				103			103		٠		103				103	

Table 5

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PDB annotation	FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-	FINGER GLI; GLI, ZINC	FINGER, COMPLEX (DNA-BINDING)	COMPLEX (DNA-BINDING	PROTEIN/DNA) FIVE-	FINGER GLI; GLI, ZINC	FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING	PROTEIN/DNA) FIVE-	FINGER GLI; GLI, ZINC	FINGER, COMPLEX (DNA-	BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING	TROIBINIDINA) FIVE-	FINGER GLI; GLI, ZINC	FINDER, COMPLEY (DIVA-	COMPLEX COA-BINDING	PROTEIN/DNA) FIVE-	FINGER GLI: GLI, ZINC	FINGER, COMPLEX (DNA-	BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING	PROTEIN/DNA) FIVE.	FINGER GLI; GLI, ZINC	FINGER, COMPLEX (DNA-	BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING	TATE CALCULATION I	FINGER COMPLEX (DNA.	BINDING PROTEIN/DNA)		CONTRACTILE PROTEIN	
Compound		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA;	CHAIN: C, D;		ZINC FINGER PROTEIN	GLII; CHAIN: A; DNA;	CHAIN: C, D;		ZINC FINGER PROTEIN	GLII; CHAIN: A; DNA;	CHAIN: C, D;			ZINC FINGER PROTEIN	GLII; CHAIN: A; DINA;	CHAIN: C, D;		ZINC KINGER PROTEIN	GLII: CHAIN: A: DNA:	CHAIN: C. D.			ZINC FINGER PROTEIN	GLII; CHAIN: A; DNA;	CHAIN: C, D;			ZINC FINGER PROTEIN	CELLI, CITOLIV. D. CIVA,	CHAIN: C. D.			TROPOMYOSIN; CHAIN: A,	
SEQ FOLD score					·										-				,						,									
PMF score		1.00			1.00				1.00					1.00				0 08	9				0.41					0.00					-0.20	
Verify score		0.50			0.38				0.32					0.49				-0.03	3				-0.01					-0.72	•				0.09	
Psi Blast		3.4e-33			le-32				1.2e-52					I.7e-33				6.88-30	0.00.0				3.4e-21					8.5e-29					1.2e-10	
END AA		432			9				464					88				408	?				504					221					442	
START		304			332				352					360				388	9				416					2					216	
CHAIN		٧			Ą				A					4				V					A					¥.	٠.				Ą	•
PDB DD		2gli			2gli				2gli					2gli				20li	7.Eu				2gli					2gli					lclg	
SEQ ID NO:		103			103				103					103				103	3				103					103		•			104	

PDB annotation	TROPOMYOSIN COILED- COIL ALPHA-HELICAL, CONTRACTILE PROTEIN	CONTRACTILE PROTEIN TROPOMYOSIN COLLED- COLL ALPHA-HELICAL, CONTRACTILE PROTEIN	CONTRACTILE PROTEIN TROPOMYOSIN COLLED- COIL ALPHA-HELICAL, CONTRACTILE PROTEIN	COMPLEX (NUCLEOCAPSID PROTEIN/RNA) NUCLEOCAPSID PROTEIN, COMPLEX (NUCLEOCAPSID PROTEIN/RNA), 2 STEM-LOOP RNA-	₽	TRANSFERASE MRNA PROCESSING, TRANSFERASE, TRANSCRIPTION, RNA- BINDING, 2 PHOSPHORYLATION, NUCLEAR PROTEIN, ALTERNATIVE SPLICING 3 II HELICAL TURN MOTIF, NUCLEOTIDYL TRANSFERASE CATALYTIC	TRANSFERASE MRNA IU PROCESSING, IL TRANSFERASE, IL
Compound	в,с, D	TROPOMYOSIN; CHAIN: A, B, C, D	TROPOMYOSIN; CHAIN: A, B, C, D	NUCLEOCAPSID PROTEIN; CHAIN: A; SL3 STEM- LOOP RNA; CHAIN: B;	NUCLEOCAPSID PROTEIN HIV-1 NUCLEOCAPSID PROTEIN (MN ISOLATE) (NMR, 20 STRUCTURES) 1AAF 3	POLYMERASE; CHAIN: A;	POLY(A) POLYMERASE; CHAIN: A;
SEQ FOLD score		·		·			
PMF		-0.19	-0.20	0.17	0.12	0.53	0.41
Verify score		0.09	0.02	-0.33	-0.53	-0.04	0.01
Psi Blast		3.4e-11	3.46-19	3.6e-07	2.4e-07	6-05	2.4e-07
END		503	503	1383	1383	542	1147
START AA		261	260	1347	1347	343	966
CHAIN		Y	A	•		Ą	· A
PDB UD		lclg	lclg	lalt	laaf	155a	lfSa
SEQ ID NO:		104	105	107	107	107	107

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PDB annotation	TRANSCRIPTION, RNA- BINDING, 2 PHOSPHORYLATION, NUCLEAR PROTEIN, ALTERNATIVE SPLICING 3 HELICAL TURN MOTIF, NUCLEOTIDYL TRANSFERASE CATALYTIC DOMAIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN						1000	ا ممر			Lines.	COMPLEX (ZINC FINGER/DNA) ZINC FINGER (FINGER	INTERACTION, PROTEIN	STRUCTURE, COMPLEX (ZINC FINGER/DNA) 71
Compound		OGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B,	TE ANSCRIPTION	REGULATION YEAST TRANSCRIPTION FACTOR ADRI (RESIDUES 102 - 130)	TERM 3 (AMINO	DOMAIN (NMR, 10	STRUCTURES) 1ARD 4	DNA-BINDING PROTEIN	HUMAN ENHANCER- RINDING PROTEIN MRP.1	MUTANT WITH CYS 11	ABIT (C11 ABIT) (NIMB. 60	STRUCTURES) 1BBO 4	DNA; CHAIN: A, B, D, B; CONSENSUS ZINC FINGER PROTEIN: CHAIN: C B G:		
SEQ FOLD score															
PMF score		0.66	070	8				0.18					1.00		
Verify		0.40	0.17	4.				0.13					0.27		•
Psi Blast		2.4e-27	2 40.10	2.46-10				1.2e-11			·		6.8e-47	·	
A A		312	797	/87				287					339		
START AA		257	250	â				257					262		
CHAIN		4											၁		
PDB ID		lalh	10.0	Lard				1bbo					Imey		
SEQ ID NO:	·	108	108	208	,			108					108		

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PDB annotation	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA	INTERACTION, PROTEIN	DESIGN, 2 CKYSTAL STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER,	PROTEIN-DNA	DESIGN, 2 CRYSTAL	STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER,	PROTEIN-DNA	DESIGN 2 CRYSTAL	STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER,	PROTEIN-DNA	DESIGN 2 CRYSTAI	STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, [I]	PROTEIN-DNA	DESIGN, 2 CRYSTAL	STRUCTURE, COMPLEX	(ZINC FINGEROUNA)		Erai E	#15
Compound	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN: CHAIN: C, F, G;		٠		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;			DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;				DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;	•			DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;			TRANSCRIPTION	REGULATION YEAST	TRANSCRIPTION FACTOR ADRI (RESIDUES 130 - 159)	
SEQ FOLD score					116.59							è.														
PMF score	1.00								1.00					68.0					0.16				0.55	}		
Verify	0.36								0.42					0.28					-0.08				-0.05			
Psi Blast	1.2e-50				1.7e-51				1.7e-51					3.4e-49					1.7e-34				0.00096			
END AA	367				368				395					424					429		•		285			
START AA	286				286				314					342		-			370				259			
CHAIN	υ <u>.</u>				D				U					၁					ນ							
PDB ED	lmey				Imey				1mey					1mey					1mey	•			lnaa		_	
SEQ ID NO:	108				801				108					801					108				108			

					
PDB annotation		ZINC FINGER TRANSCRIPTION FACTOR SP1; ZINC FINGER, TRANSCRIPTION ACTIVATION, SP1	COMPLEX (TRANSCRIPTION REGULATION/DNA) TFIILA; 5S GENE; NMR, TFIILA, PROTEIN, DNA, TRANSCRIPTION PACTOR, 5S RNA 2 GENE, DNA BINDING PROTEIN, ZINC FINGER, COMPLEX 3 (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION— REGULATIONDNA) COMPLEX (TRANSCRIPTION) REGULATIONDNA), RNA POLYMERASE III, 2
Compound	IPAA 3 (PAPA - CARBOXY TERMINAL ZINC FINGER DOMAIN) MUTANT WITH IPAA 4 PRO 131 REPLACED BY ALA, PRO 133 REPLACED BY ALA, CYS 140 IPAA 5 REPLACED BY ALA (P131A,P133A,C140A) (NMR, 10 STRUCTURES) IPAA 6	SPIPS; CHAIN: NULL;	TRANSCRIPTION FACTOR IIIA; CHAIN: A; 5S RNA GENE; CHAIN: E, F;	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;
SEQ FOLD score	·			94.31	
PMF		0.46	0.36	,	0.99
Verify score		0.30	0.32		0.05
Psi Blast		1.1e-09	7.2e-15	1.2e-35	1.2e-35
A END		287	314	428	404
START		259	257	259	262
CHAIN			A	¥	¥
EDB CE		1sp2	1473	1tf6	1116
SEQ ID NO:		108	108	108	108

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PDB annotation	TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION) REGULATION/DNA) (COMPLEX (TRANSCRIPTION) REGULATION/DNA) YING- YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELMENT, YY1, ZINC 2 FINGER PROTEIN, DNA- PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION) REGULATION/DNA)	COMPLEX (TRANSCRIPTION)
Compound		TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YY1; CHAIN: C; ADENO-
SEQ FOLD score				90.80		
PMF score		0.62	1.00		0.53	90.0
Verify		-0.10	0.30		-0.06	-0.16
Psi Blast		1.7e-34	1 c- 32	8.5e-33	8.5e-33	1.7e-24
END		426	367	396	424	429
START		287	266	284	322	350
CHAIN		A	U	IO	U	ပ
EDB EDB		1466	Inbd	Iubd	lubd	1ubd
SEQ ID		108	108	108	108	108

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PDB annotation	REGULATION/DNA) YING- YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA- PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (DNA-BINDING PROTBINDNA) FIVE- FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE- FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTBIN/DNA) FIVE- FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE- FINGER GLI, GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	MYOSIN MYOSIN MOTOR	MYOSIN MYOSIN MOTOR
Compound	ASSOCIATED VIRUS P5 INTTIATOR ELEMENT DNA; CHAIN: A, B;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	MYOSIN HEAVY CHAIN; CHAIN; A; MYOSIN REGULATORY LIGHT CHAIN; CHAIN: Y; MYOSIN ESSENTIAL LIGHT CHAIN; CHAIN: Z;	MYOSIN HEAVY CHAIN; CHAIN: A; MYOSIN REGULATORY LIGHT CHAIN; CHAIN: Y;;
SEQ FOLD score			97.44			483.37	
PMF score		1.00		1:00	0.54		1.00
Verify		0.22		0.29	0.26		0.50
Psi Blast		4.8e-66	4.8e-66	6.8e-33	8.56-31	o	0
END		966	966	394	424	784	789
START AA		259		266	294	-	6
CHAIN		A	Ą	¥	V	∢	· V
PDB ID		2gli	2gli	2gli	2gli	1b7t	1b7t
SEQ ID NO:		108	801	108	108	109	109

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PDB annotation		MUSCLE PROTEIN MDE; MUSCLE PROTEIN	MUSCLE PROTEIN MDE; MUSCLE PROTEIN	MUSCLE PROTEIN MUSCLE PROTEIN	MUSCLE PROTEIN MUSCLE PROTEIN	CONTRACTILE PROTEIN MYOSIN MOTOR, CONFORMATIONAL CHANGES	CONTRACTILE PROTEIN MYOSIN, DICTYOSTELIUM,	MOTOR, MANT, ATPASE, ACTIN-BINDING, 2 COILED COIL	CONTRACTILE PROTEIN	MOTOR, MANT, ATPASE, ACTIN-BINDING, 2 COILED COIL	CONTRACTILE PROTEIN	ATPASE, MYOSIN, COILED COIL ACTIN-BINDING, ATP.	BINDING, 2 HEPTAD	REPEAT PATTERN, METHYLATION,	ALKYLATION, 3	PHOSPHORYLATION, CONTRACTILE PROTEIN	CONTRACTILE PROTEIN ATPASE, MYOSIN, COILED	COLL, ACTIN-BINDING, ATP-BINDING, 2 HEPTAD	REPEAT PATTERN, METHYLATION,
Compound	MYOSIN ESSENTIAL LIGHT CHAIN; CHAIN: Z	MYOSIN; CHAIN: A, B, C, D, E, F, G, H;	MYOSIN; CHAIN: A, B, C, D, E, F, G, H;	MYOSIN; CHAIN: A, B, C, D, E, F;	MYOSIN; CHAIN: A, B, C, D, E, F,	MYOSIN HEAD; CHAIN: A; MYOSIN HEAD; CHAIN: Y; MYOSIN HEAD; CHAIN: Z;	MYOSIN; CHAIN: NULL;		MYOSIN; CHAIN: NULL;		MYOSIN; CHAIN: NULL;		•		•		MYOSIN; CHAIN: NULL;		
SEQ FOLD score			526.30		489.18	·	496.50				425.46								
PMF score		1.00		1.00		1.00		. :	1.00					. *			1.00		
Verify score		0.61		0.65		0.33			0.48								0.40	·	
Psi Blast		0	0	0	0	0	0		0		0			····			0		
END AA		740	740	710	710	789	710		710		639						639		
START AA		1		1		O	-		9		1						3		
CHAIN		¥	A	A	Ą	V									-				
PDB UD		1br1	1br1	1br2	1br2.	1dfk	llvk		livk		Imnd						lmnd		
SEQ ID NO:		109	109	109	109	109	109		109		109						109		

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PDB annotation	ALKYLATION, 3 PHOSPHORYLATION, CONTRACTILE PROTEIN c	MUSCLE PROTEIN MUSCLE PROTEIN, MYOSIN SUBFRAGMENT-1, MYOSIN	MUSCLE PROTEIN MUSCLE PROTEIN, MYOSIN SUBFRAGMENT-1, MYOSIN HEAD A MOTOR PROTEIN	HEAD, 2 MOLON FROIDEN	KINASE KINASE, SIGNAL TRANSDUCTION, CALCIUM/CALMODULIN	KINASE KINASE, SIGNAL TRANSDUCTION, CALCIUM/CALMODULIN	TRANSFERASE, TRANSFERASE, SERINE/THREONINE. PROTEIN KINASE, CASEIN KINASE, 2 SER/THR KINASE		
Compound		MYOSIN; CHAIN: A, B, C;	MYOSIN; CHAIN: A, B, C;		CALCIUM/CALMODULIN- DEPENDENT PROTEIN KINASE; CHAIN: NULL;	CALCTUM/CALMODULIN- DEPENDENT PROTEIN KINASE; CHAIN: NULL;	PROTEIN KINASE CK2/ALPHA-SUBUNIT; CHAIN: NULL;	TRANSFERASE(PHOSPHO TRANSFERASE) \$C-/AMP\$- DEPENDENT PROTEIN KINASE (B.C.2.7.1.37) (\$C/APK\$) 1APM 3 (\$C/APK\$) 1APM 3 ALPHA ISOENZYME MUTANT WITH SER 139 1APM 4 REPLACED BY ALA (\$1394\$) COMPLEX WITH THE PEPTIDE 1APM 5 INHBITOR PKI(5-24) AND THE DETERGENT MEGA-8 1APM 6	TRANSFERASE(PHOSPHO TRANSFERASE) \$C-/AMP\$-
SEQ FOLD score		392.48			142.86		110.78		168.62
PMF			1.00			1.00		1.00	
Verify score			0.58			0.39		0.54	
Psi Blast		0	0		1.7e-89	1.7e-89	3.6e-56	0	0
END AA		771	743		318	300	321	311	326
START AA		1	9		10	11		17	2
CHAIN	·	Ą	. V				·	m	ш
PDB ED		2mys	2mys		1a06	1a06	1a60	lapm	lapm
SEQ ID NO:		601	109		. 110	110 ·	110	110	110

		· · · · · · · · · · · · · · · · · · ·			- 11122)
PDB annotation		PROTEIN KINASE CDK2; PROTEIN KINASE, CELL CYCLE, PHOSPHORYLATION, STAUROSPORINE, 2 CELL DIVISION, MITOSIS, INHIBITION	PROTEIN KINASE CDK2; PROTEIN KINASE, CELL CYCLE, PHOSPHORYLATION, STAUROSPORINE, 2 CELL DIVISION-MITOSIS, INHIBITION			PHOSPHOTRANSFERASE
Compound	DEPENDENT PROTEIN KINASE (B.C.2.7.1.37) (\$C'APK\$) 1APM 3 (CATAL YTIC SUBUNIT) ALPHA ISOBNZYME MUTANT WITH SER 139 1APM 4 REPLACED BY ALA (\$139A\$) COMPLEX WITH THE PEPTIDE 1APM 5 INHIBITOR PKI(5-24) AND THE DETERGENT MEGA-8 1APM 6	CYCLIN-DEPENDENT PROTEIN KINASE 2; CHAIN: NULL;	CYCLIN-DEPENDENT PROTEIN KINASE 2; CHAIN: NULL;	PHOSPHOTRANSFERASE CAMP-DEPENDENT PROTEIN KINASE CATALYTIC SUBUNIT ICMK 3 (E.C.2.7.1.37)	PHOSPHOTRANSFERASE CAMP-DEPENDENT PROTEIN KINASE CATALYTIC SUBUNIT 1CMK 3 (E.C.2.7.1.37) 1CMK 4	CASEIN KINASE-1; 1CSN 4
SEQ FOLD score		112.67			163.96	
PMF score			1.00	1.00		1.00
Verify score		-1	0.34	0.64		0.49
Psi Blast		3.6e-68	3.66-68	0	o ·	2.4e-66
END		285	271	311	329	274
START AA		17	20 .	91	2	18
CHAIN				团	m m	
PDB ID		1aq1	1aq1	1cmk	1cmk	1csn
SEQ ID NO:	·	110	110	110	110	110

Table 5

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PDB annotation			PHOSPHOTRANSFERASE FGFR IK, FIBROBLAST GROWTH FACTOR RECEPTOR 1; TRANSFERASE, TYROSINE-PROTEIN KINASE, ATP-BINDING, 2 PHOSPHORYLATION, RECEPTOR, PHOSPHOTRANSFERASE	PHOSPHOTRANSFERASE FGRRIK, FIBROBLAST GROWTH FACTOR RECEPTOR 1; TRANSFERASE, TYROSINE- PROTEIN KINASE, ATP- BINDING, 2 PHOSPHORYLATION, RECEPTOR, RECEPTOR,	PROTEIN KINASE CDK2; TRANSFERASE, SERINETHREONÎNE PROTEIN KINASE, ATP- BINDING, 2 CELL CYCLE,
Compound	TRANSFERASE(PHOSPHO TRANSFERASE) CAMP- DEPENDENT PROTEIN KINASE (B.C.2.7.1.37) (CAPK) 1CTP 3 (CATALYTIC SUBUNIT)	TRANSFERASE(PHOSPHO TRANSFERASE) CAMP- DEPENDENT PROTEIN KINASE (B.C.2.7.1.37) (CAPK) 1CTP 3 (CATAL YTIC SUBUNIT)	FGF RECEPTOR 1; CHAIN: A, B;	FGF RECEPTOR 1; CHAIN: A, B;	HUMAN CYCLIN- DEPENDENT KINASE 2; CHAIN: NULL;
SEQ FOLD score	·	171.30	119.24	138.09	122.03
PMF score	00.1				
Verify score	0.77				
Psi Blast	0	0	3.4e-34	4.8e-47	8.5e-62
END	304	313	277	276	285
START AA	16	2	. 7	-	17
CHAIN	щ	E	V .	m	·
PDB ID	1сф	1сф	1fgk	1fgk	Ihci
SEQ ID	110	110	110	011	110

Table 5

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PDB annotation	CELL DIVISION, MITOSIS, PHOSPHORYLATION	TRANSFERASE INK3; TRANSFERASE, INK3 MAP KTNASF	SERINE/THREONINE PROTEIN 2 KINASE	KINASE KINASE, TWITCHIN, INTRASTERIC REGULATION	KINASE KINASE, TWITCHIN, INTRASTERIC REGULATION	KINASE KINASE, TWITCHIN, INTRASTERIC REGULATION	KINASE KINASE, TWITCHIN, INTRASTERIC REGULATION	TRANSFERASE MITOGEN	KINASE; TRANSFERASE,	MAP KINASE,	SERINE/THREONINE- PROTEIN KINASE, 2 P38	KINASE RABBIT MUSCLE PHOSPHORYLASE KINASE;	GLYCOGEN METABOLISM,	TRANSFERASE,	PROTEIN, 2 KINASE, ATP-	BINDING, CALMODULIN-	BINDING TO THE STREET THE	PHOSPHORYT ASE KINASE:	GLYCOGEN METABOLISM,	TRANSFERASE,	SERINE/THREONINE-	PROTEIN, 2 KINASE, ATP-	BINDING	SERINE KINASE SERINE KINASE, TITIN, MUSCLE,
Compound		C-IUN N-TERMINAL KINASE; CHAIN: NULL;		TWITCHIN; CHAIN: NULL;	TWITCHIN; CHAIN: A, B;	TWITCHIN; CHAIN: A, B;	TWITCHIN; CHAIN: A, B;	MAP KINASE P38; CHAIN:	NOTE,			PHOSPHORYLASE KINASE; CHAIN: NULL;					TO A MANAGEMENT OF THE	FHOSPHOK YLASE KTNASE: CHATN: NITL:						TITIN; CHAIN: A, B;
SEQ FOLD score		127.21			139.53			119.80									0000	170.32						124.66
PMF				1.00		1.00	1.00					1.00												
Verify				0.48		0.39	0.36					0.77												
Psi Blast		3.4e-48		1.7e-73	1.2e-92	1.2e-73	1.2e-92	2.4e-63				5.1e-88					3	3.1e-88						1.2e-60
END A		366		275	344	274	336	313				272						275						331
START		4		9	3	9	7	5				18						<u> </u>					,	16
CHAIN					Ą	A	A																	A
PDB UD		1jnk		1koa	Ikob	1kob	1kob	1p38				1phk						1phk		_				1tki
SEQ ID NO:		110		110	110	110	110	110				110						011						110

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					_	-			_	•				_				-			7	4	D. 1		D	-		1	1	
PDB annotation	AUTOINHIBITION	TRANSFERASE MITOGEN ACTIVATED PROTEIN KINASH MAP 2 HRK2:	TRANSFERASE,	SERINE/THREONINE	FROTEIN KINASE, MAP KINASE, 2 ERK2		TOXIN BINDING PROTEIN	PROPELLER AND	ALPHA/BETA FOLD	TRANSCRIPTION INHIBITOR BETA-PROPELLER		TRANSCRIPTION INHIBITOR	BETA-PROPELLER	COMPLEX (GTP-	BINDING/TRANSDUCER)	BETAI, TRANSDUCIN BETA	SUBUNIT; GAMMAI,	TRANSDUCIN GAMMA	SUBUNIT; COMPLEX (GIF-	PROTEIN HETEROTRIMER 2	SIGNAL TRANSDUCTION	COMPLEX (GTP-	BINDING/TRANSDUCER)	BEI'AI, TRANSDUCIN BEI'A	SUBUNIT; GAMMAI,	FIRANSDUCIN GAMMA	RINDING/TRANSDITCER) G	PROTEIN HETEROTRINER 2	SIGNAL TRANSDUCTION	COMPLEX (GTP- BINDING/TRANSDUCER)
Compound		EXTRACELLULAR REGULATED KINASE 2; CHAIN: MITT:			•		TOLB PROTEIN; CHAIN: A;			TRANSCRIPTIONAL REPRESSOR TUP1: CHAIN:	A, B, C;	TRANSCRIPTIONAL	REPRESSOR TUP1; CHAIN: A. B. C:	GT-ALPHA/GI-ALPHA	CHIMERA; CHAIN: A; GT-	BETA; CHAIN: B; GT-	GAMMA; CHAIN: G;	•				GT-ALPHA/GI-ALPHA	CHIMERA; CHAIN: A; GT.	BEI'A; CHAIN: B; GI-	GAMMA; CHAIN: G;		· ·			GT-ALPHA/GI-ALPHA CHIMERA; CHAIN: A; GT-
SEQ FOLD score		120.67			-									167.73												٠				
PMF		·					0.31			1.00		1.00				•						86.0								1.00
Verify							0.20			0.49		0.94							, .			0.48								0.89
Psi Blast		1.2e-47					0.00017			3.4e-70		8.5e-73		1.7e-83								1.7e-47		-						1.7e-83
AA END		370					400			473		514		470		-						377			-	-,		<u>-</u>		470
START AA		4					9/1			146		224		114								139								156
CHAIN							Ą			∢		A		В			-					В								æ
PDB CI		3erk					lorz			lerj		lerj		1got								1 got								1got
SEQ ID NO:		110					111			111		111		1111								111								111

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				71			SOLE	'Uleb	
PDB annotation	BETAI, TRANSDUCIN BETA SUBUNIT; GAMMAI, TRANSDUCIN GAMMA SUBUNIT; COMPLEX (GTP- BINDING/TRANSDUCER), G PROTEIN, HETEROTRIMER 2 SIGNAL TRANSDUCTION	COMPLEX (GTP. BINDING/TRANSDUCER) BETA1, TRANSDUCIN BETA SUBUNIT; GAMMA1, TRANSDUCIN GAMMA SUBUNIT; COMPLEX (GTP- BINDING/TRANSDUCER), G PROTEIN, HETEROTRIMER 2 SIGNAL TRANSDUCTION	REPLICATION DNA NUCLEOTIDE EXCISION REPAIR, UVRABC, HELICASE, 2 HYPERTHERMOSTABLE	HYDROI ASE ITVER:	MULTIDOMAIN PROTEIN	GENE REGULATION APO PROTEIN	TRANSLATION YEAST INITIATION FACTOR 4A, EIF4A; HELICASE, INITIATION FACTOR 4A, DEAD-BOX PROTEIN	TRANSLATION EUKARYOTIC INITIATION FACTOR 4A; IF4A, HELICASE, DEAD-BOX PROTEIN	TRANSLATION
Compound	BETA; CHAIN: B; GT- GAMMA; CHAIN: G;	GT-ALPHA/GI-ALPHA CHIMERA; CHAIN: A; GT- BETA; CHAIN: B; GT- GAMMA; CHAIN: G;	DNA NUCLEOTIDE EXCISION REPAIR ENZYME UVRB; CHAIN: A;	EVCTATIOT BACE ABC	EXCINUCLEASE ABC SUBUNIT B; CHAIN: A;	EXCINUCLEASE UVRABC COMPONENT UVRB; CHAIN: A;	EUKARYOTIC INITIATION FACTOR 4A; CHAIN: A;	YEAST INITIATION FACTOR 4A; CHAIN: A, B;	YEAST INITIATION
SEQ FOLD score									
PMF		1.00	0.69	20.0	0.25	0.35	-0.05	0.06	0.04
Verify		0.79	0.58	0 50	0.52	0.07	0.10	-0.10	-0.31
Psi Blast		5.1e-68	3.40-10	3.45.10	3.4e-10	1e-14	5.1e-36	0.0024	1.2e-91
END A	·	511	431	421	431	468	445	120	445
START		219	301	301	301	301	285	7	56
CHAIN		В	Ą		A	Ą	∢	В	В
PDB		1got	1040	1.40	1d2m	149x	1fûk	Ifuu	1fm
SEQ ID		111	112		112	112	112	112	112

Table 5

			·	 		4 4		di alla fizza fizza	- 1225
PDB annotation	EUKARYOTIC INITIATION FACTOR 4A; IF4A, HELICASE, DEAD-BOX PROTEIN	CHAPERONE/STRUCTURAL PROTEIN CHAPERONE ADHESIN DONOR STRAND COMPLEMENTATION, 2 CHAPERONE/STRUCTURAL PROTEIN	CHAPERONE/STRUCTURAL PROTEIN CHAPERONE ADHESIN DONOR STRAND COMPLEMENTATION, 2 CHAPERONE/STRUCTURAL PROTEIN	HYDROLASB HYDROLASE, DEPHOSPHORYLATION	HYDROLASE PTP1B; HYDROLASE, PHOSPHORYLATION, LIGAND, INHIBITOR	HYDROLASE C2 DOMAIN, PHOSPHOTIDYLINOSITOL, PHOSPHOTASE, HYDROLASE	HYDROLASE PROTEIN- TYROSINE PHOSPHATASE; HYDROLASE, PROTBIN TYROSINE PHOSPHATASE, CATALYTIC DOMAIN, 2 WPD LOOP, SH2 DOMAIN	HYDROLASE DUAL SPECIFICITY PHOSPHATASE, MAP KINASE HYDROLASE	HYDROLASE DUAL
Compound	FACTOR 4A; CHAIN: A, B;	PAPD-LIKE CHAPERONE FIMC; CHAIN: A, C, B, G, I, K, M, O; MANNOSE- SPECIFIC ADHESIN FIMH; CHAIN: B, D, F, H, I, L, N, P;	PAPD-LIKE CHAPERONE FIMC; CHAIN: A, C, B, G, I, K, M, O; MANNOSE- SPECIFIC ADHESIN FIMH; CHAIN: B, D, F, H, I, L, N, P;	PROTEIN TYROSINE PHOSPHATASE 1B; CHAIN: NULL;	PROTEIN-TYROSINE PHOSPHATASE 1B; CHAIN: A;	PHOSPHOINOSITIDE PHOSPHOTASE PTEN; CHAIN: A;	SHP-1; CHAIN: NULL;	PYST1; CHAIN: NULL;	PYST1; CHAIN: NULL;
SEQ FOLD score					·				
PMIF		-0.20	-0.20	0.03	-0.09	0.95	0.01	0.71	0.48
Verify		0.14	0.29	0.25	0.20	0.38	0.18	0.14	-0.04
Psi Blast		1.2e-08	9.6e-10	1.7e-35	1.7e-38	3.4e-20	8.5e-40	2.4e-20	1.7e-12
END A		1347	1307	293	293	305	301	296	296
START		1227	1241	68	68	122	68	159	185
CHAIN		æ	В		Ą	A			
PDB		1qun	lqun	1a5y	1c83	1d5r	1gwz	1mkp	1mkp
SEQ ID		112	112	114	114	114	114	114	114

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			_						
PDB annotation	SPECIFICITY PHOSPHATASE, MAP KINASE HYDROLASE	RECEPTOR DI; RECEPTOR, PHOSPHATASE, SIGNAL TRANSDUCTION, ADHESION, 2 HYDROLASE	HYDROLASE VHR; HYDROLASE, PROTEIN DUAL-SPECIFICITY PHOSPHATASE	HYDROLASE VHR; HYDROLASE, PROTEIN DUAL-SPECIFICITY PHOSPHATASE	HYDROLASE D1; HYDROLASE, SIGNAL TRANSDUCTION, RECEPTOR, GLYCOPROTEIN, 2 PHOSPHORYLATION, SIGNAL	HYDROLASE YOP51, YOP2B, PASTEURELLA X, PTP-ASE, PROTEIN TYROSINE PHOSPHATASE, HYDROLASE	HYDROLASE YOP51, YOP2B, PASTEURELLA X, PTP-ASB, E PROTEIN TYROSINE PHOSPHATASE, HYDROLASE	TYROSINE PHOSPHATASE SYP, SHPTP-2; TYROSINE PHOSPHATASE, INSULIN SIGNALING, SH2 PROTEIN	HYDROLASE SUMO HYDROLASE, UBIQUITIN-
Compound		RECEPTOR PROTEIN TYROSINE PHOSPHATASE MU; CHAIN: A, B;	HUMAN VHI-RELATED DUAL-SPECIFICITY PHOSPHATASE CHAIN: A, B:	HUMAN VHI-RELATED DUAL-SPECIFICITY PHOSPHATASE CHAIN: A, B;	RECEPTOR PROTBIN TYROSINB PHOSPHATASE ALPHA; CHAIN: A, B;	YERSINIA PROTEIN TYROSINE PHOSPHATASE; CHAIN: NULL;	YERSINIA PROTEIN TYROSINE PHOSPHATASE; CHAIN: NULL;	SHP-2; CHAIN: A, B;	ULPI PROTEASE; CHAIN: A; UBITQUTIN-LIKE
SEQ FOLD score									
PMF		0.09	0.99	0.40	60:0	0.00	0.39	-0.12	1.00
Verify		-0.10	0.49	0.38	0.31	-0.05	-0.07	0.17	0.51
Psi Blast		1.5e-43	2.4e-20	6.8e-11	6.8e-46	6.8e-12	3.6e-05	1.4e-48	5.1e-41
EZ Y		298	296	301	298	289	294		288
START		84	124	160	29	115	205	59	364
CHAIN		¥	Ą	A	¥			V	A
PDB		lrpm	lvhr	lvhr	1yfo	lytn	lytn	2shp	leuv
SEQ ID		114	114	114	114	114	114	114	115

Fable:

					_				Harrier Harris
PDB annotation	LIKE PROTEASE 1, SMT3 HYDROLASE 2 DESUMOYLATING ENZYME, CYSTEINE PROTEASE, SUMO PROCESSING 3 ENZYME, SMT3 PROCESSING ENZYME, NABH4, THIOHEMIACETAL, 4 COVALENT PROTEASE ADDUCT		ENDOCYTOSIS/EXOCYTOSI S SYNAPTOTAGMIN ASSOCIATED 35 KDA PROTEIN, P35A, THREE HELIX BUNDLE	ENDOCYTOSIS/EXOCYTOSI S SYNAPTOTAGMIN ASSOCIATED 35 KDA PROTEIN, P35A, THREE HELIX BUNDLE	ENDOCYTOSIS/EXOCYTOSI	S SYNAPTOTAGMIN ASSOCIATED 35 KDA PROTEIN, P35A, THREE HELIX BUNDLE	ENDOCYTOSIS/EXOCYTOSI S SYNAPTOTAGMIN ASSOCIATED 35 KDA PROTEIN, P35A, THREE HELIX BUNDLE	ENDOCYTOSIS/EXOCYTOSI S SYNAPTOTAGMIN ASSOCIATED 35 KDA PROTEIN, P35A, THREE HELIX BUNDLE	ENDOCYTOSIS/EXOCYTOSI S SYNAPTOTAGMIN
Compound	PROTEIN SMT3; CHAIN: B;		SYNTAXIN-1A; CHAIN: A, B, C;	SYNTAXIN-1A; CHAIN: A, B, C;	SYNTAXIN-1A; CHAIN: A,	ပံ ရ	SYNTAXIN-1A; CHAIN: A, B, C;	SYNTAXIN-1A; CHAIN: A, B, C;	SYNTAXIN-1A; CHAIN: A, B, C;
SEQ FOLD score								-	
PMF			-0.17	-0.17	-0.06		-0.18	-0.19	-0.18
Verify			0.29	0.28	0.15		0.15	0.25	0.11
Psi Blast			6e-11	4.8e-09	1.2e-11		2.4e-11	1.16-10	8.4e-10
A END		j	265	293	327		425	499	151
START			145	174	206		307	378	52
CHAIN			¥	¥	¥		V	∢	A
PDB U			lez3	lez3	lez3		1ez3	lez3	lez3
SEQ ID NO:			118	118	118		118		118

				r———			700. () () (100° 3° 11	() - () - () - () () () () () (
PDB annotation	ASSOCIATED 35 KDA PROTEIN, P35A, THREE HELIX BUNDLE	ENDOCYTOSIS/EXOCYTOSI S SYNAPTOTAGMIN ASSOCIATED 35 KDA PROTEIN, P35A, THREE HELIX BUNDLE	CONTRACTILE PROTEIN TRIPLE-HELIX COLLED COLL, CONTRACTILE PROTEIN	TRANSCRIPTION REGULATION SIGMA70; RNA POLYMERASE SIGMA PACTOR, TRANSCRIPTION REGULATION		G day	METAL TRANSPORT INHIBITOR/RECEPTOR HFB; (HFB, HEREDITARY HEMOCHROMATOSIS, MHC CLASS I, TRANSFERRIN 2 RECEPTOR	HYDROLASE SGAP; DOUBLE-ZINC METALLOPROTEINAZE, CALCIUM ACTIVATION, PROTEIN- 2 INHIBITOR
Compound		SYNTAXIN-1A; CHAIN: A, B, C;	HUMAN SKELETAL MUSCLE ALPHA-ACTININ 2; CHAIN: A;	RNA POLYMERASE PRIMARY SIGMA FACTOR; CHAIN: NULL;	HYDROLASE(AMINOPEPT IDASE) AMINOPEPTIDASE (AEROMONAS PROTEOLYTICA) (E.C.3.4.11.10) 1AMP 3	HYDROLASE(AMINOPEPT IDASE) AMINOPEPTIDASE (AEROMONAS PROTEOLYTICA) (E.C.3.4.11.10) 1AMP 3	HEMOCHROMATOSIS PROTEIN; CHAIN: A, D, G; BETA-2- MICROGLOBULIN; CHAIN: B, E, H; TRANSFERRIN RECEPTOR; CHAIN: C, F, I;	AMINOPEPTIDASE; CHAIN: A;
SEQ FOLD score				•				
PMF score		-0.18	-0.12	0.06	0.43	0.43	0.83	0.64
Verify score		0.08	0.02	-0.24	0.04	0.11	0.02	-0.00
Psi Blast		8.4e-13	6e-16	1.2e-09	1.2e-18	5.1e-30	16-46	1.7e-28
END		224	116	251	356	468	292	463
START AA		93	736	52	193	211	224	211
CHAIN		Α .	Ą		·		ပ	¥
PDB ID		lez3	nnbj	lsig .	lamp	lamp	1de4	1949
SEQ ID NO:		118	118	118	120	120	120	120

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PDB annotation	COMPLEX		STRUCTURAL PROTEIN RETINAL S-ANTIGEN, 48 KD	PROTEIN; VISUAL ARRESTIN	DESENSITISATION OF THE	VISUAL TRANSDUCTION 2	CASCADE, BINDING TO	ACTICATED AND	PHOSPHORYLATED RHODOPSIN	STRUCTURAL PROTEIN RETINAL S-ANTIGEN 48 KD	PROTEIN; VISUAL	ARRESTIN,	DESENSITISATION OF THE VISITAL TRANSDICTION 2	CASCADE, BINDING TO	ACTICATED AND	PHOSPHORYLATED RHONOPSIN	STRUCTURAL PROTEIN	RETINAL S-ANTIGEN, 48 KD	PROTEIN; VISUAL	PERINGHIS ATTON OF THE	VISUAL TRANSDUCTION 2	CASCADE, BINDING TO	ACTICATED AND	PHOSPHORYLATED	KHODOPSIN	PROTEASE PROSOME,	MULTICATALYTIC	PROTEASE, MCP, MACROPAIN; PROTEASE,	PROTEASOME, HYDROLASE	MULTICATALYTIC F
Compound			ARRESTIN; CHAIN: A, B, C, D:			•		:		ARRESTIN; CHAIN: A, B, C,	i		,				ARRESTIN; CHAIN: A, B, C,	ű								PROTEASOME; CHAIN: A,	B, C, D, B, F, G, H, I, J, K, L,	M, N, O, P, Q		20S PROTEASOME;
SEQ FOLD score			73.18					,									71.95		•			··				71.75				55.61
PMF. score								-		0.00									,		•									
Verify								-		-0.35																		······································		
Psi Blast			1.2e-41							1.2e-41							1.2e-43									1.2e-44				1.46-43
END			358		•			•		265			•				351						•			205				205
START AA			1						•	59			:				1			-				-		12				-
CHAIN			¥							Ą							D					•				В				H
PDB ID		-	lcfl			_				lcfl		<u>.</u>					lefi								1	1pma				lryp
SEQ ID NO:			121							121							121									122				122.

Table 5

							
PDB annotation	PROTEINASE MULTICATALYTIC PROTEINASE, 20S PROTEASOME, PROTEIN 2 DEGRADATION, ANTIGEN PROCESSING, HYDROLASE,	MULTICATALYTIC PROTBINASE MULTICATALYTIC PROTBINASE, 20S PROTBINASE, PROTBIN 2 DEGRADATION, ANTIGEN PROCESSING, HYDROLASE, PROTFASS	MULTICATALYTIC PROTEINASE MULTICATALYTIC PROTEINASE, 20S PROTEASOME, PROTEIN 2 DEGRADATION, ANTIGEN PROCESSING, HYDROLASE, PROTEASE	MULTICATALYTIC PROTEINASE MULTICATALYTIC PROTEINASE, 20S PROTEINASE, 20S PROTEASOME, PROTEIN 2 DEGRADATION, ANTIGEN PROCESSING, HYDROLASE, (RIBOSOMAL PROTEIN, RIBOSOMAL PROTEIN, RIBOSOMAL PROTEIN, RRNA-BINDING	RIBOSOMAL PROTEIN RIBOSOMAL PROTEIN, RRNA-BINDING	lba
Compound	CHAIN: A, B, C, D, B, F, G, H, I, I, K, L, M, N, O, P, Q,	20S PROTEASOME; CHAIN: A, B, C, D, B, F, G, H, I, I, K, L, M, N, O, P, Q,	20S PROTEASOME; CHAIN: A, B, C, D, B, F, G, H, I, J, K, L, M, N, O, P, Q,	20S PROTEASOME; CHAIN: A, B, C, D, E, F, G, H, I, J, K, L, M, N, O, P, Q,	RIBOSOMAL PROTEIN 19; CHAIN: NULL;	RIBOSOMAL PROTEIN 1.9; CHAIN: NULL;	
SEQ FOLD score		84.34	58.38	52.75	·	54.38	
PMF score					0.82		
Verify score	:				0.38		
Psi Blast		1.5e-36	1.5e-36	3.46-48	6.8e-42	6.8e-42	
END		205	205	205	240	243	
START AA		1	н	2	94	94	
CHAIN		ſ	×	<u>.</u>			
PDB ID		lryp	Ігур	lryp	ldiv	1div	
SEQ ID NO:		122	221	122	124	124	

				,	1 Her 11	. If so the transfer	- 1 1 1	den thus that
PDB annotation	HALOPEROXIDASE BROMOPEROXIDASE L, HALOPEROXIDASE L; HALOPEROXIDASE, OXIDOREDUCTASE	HALOPEROXIDASE CHLOROPEROXIDASE A1, HALOPEROXIDASE A1; HALOPEROXIDASE, OXIDOREDUCTASE	HALOPEROXIDASE HALOPEROXIDASE F; HALOPEROXIDASE, OXIDOREDUCTASE, PROPIONATE COMPLEX	AMINOPEPTIDASE AMINOPEPTIDASE, PROLINE IMINOPEPTIDASE, SERINE PROTEASE, 2 XANTHOMONAS CAMPESTRIS	HYDROLASE HYDROLASE, HALOALKANE DEHALOGENASE, ALPHABETA-HYDROLASE	HALOPEROXIDASE HALOPEROXIDASE A2, CHLOROPEROXIDASE A2; HALOPEROXIDASE, OXIDOREDUCTASE, PEROXIDASE, ALPHABETA 2 HYDROLASE FOLD, MUTANT M99T	HYDROLASE BPHD; HYDROLASE, PCB DEGRADATION	HYDROLASE A/B HYDROLASE FOLD, DEHALOGENASE I-S BOND
Compound	CHAIN: A, B, C;	BROMOPEROXIDASE A1; CHAIN: NULL;	CHLOROPEROXIDASE F; CHAIN: NULL;	PROLINE IMINOPEPTIDASE; CHAIN: A, B;	HALOALKANE DEHALOGENASE; CHAIN: NULL;	BROMOPEROXIDASE A2; CHAIN: NULL;	2-HYDROXY-6-OXO-6- PHENYLHEXA-2,4- DIENOATE CHAÎN: A;	HALOALKANE DEHALOGENASE; 1- CHLOROHEXANE CHAIN:
SEQ FOLD score	68.67	54.47	50.66	56.93	71.01	68.50	63.30	73.30
PMF score								
Verify								
Psi Blast	1.4e-46	1.7e-40	1.7e-44	5.1e-37	1.2e-31	1.7e-38	1.4e-40	6.8e-43
END	366	366	366	367	353	365	998	368
START AA	74	11	22	59	51	27	<i>L</i> 9	65
CHAIN	V	·					¥	A
PDB U	1a88	1a8q	la8s	lazw	lb6g	lbrt	1c4x	lcqw
SEQ ID NO:	128	. 128	128	128	128	128	128	128

AA AA
74 365 1.7e-45
•
342 1.7e-06
93 368 8.5e-17
59 366 3.4e-35
·
84 654 0
•
93 650 0
232 0
189 350 3.4e-20

Table 5

PDB annotation	Zi.	COMPLEX	DNA), ZINC		TION POZ	TEIN- RACTION		ONAL 2	INC-FINGER	RAPHY, 3	JCTURE,	TIC	SINE		AC ZINC HINGER.		, PROTEIN	(STAL	COMPLEX	COMPLEX (TRANSCRIPTION)	DNA)	COMPLEX (TRANSCRIPTION)	DNA), RNA	7117	NC	INC FINGER	Z	ISCRIPTION OF	WITTATOR I
ius ACA	GLYCOPROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX	(ZINC FINGER/DNA), ZINC	PROTEIN	GENE REGULATION POZ	DOMAIN; PROTEIN- PROTEIN INTERACTION	DOMAIN,	TRANSCRIPTIONAL 2	REFRESSOR, ZING-FINGER PROTEIN X-RAY	CRYSTALLOGRAPHY, 3	PROTEIN STRUCTURE,	PROMYELOCYTIC	LEUKEMIA, GENE	NECOLATION.	COMPLEX (ZINC) FINGER.	PROTEIN-DNA	INTERACTION, PROTEIN	DESIGN, 2 CRYSTAL	STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX	REGULATION/DNA)	COMPLEX (TR	REGULATION/DNA), RNA	POLYMERASE III, 2	TRANSCRIPTION	PROTEIN	COMPLEX (TR	YANG 1: TRANSCRIPTION	INTITATION, INITIATOR
Compound		OGSR ZINC FINGER PEPTIDE; CHAIN: A;	DUPLEX OF ICOMFICE POPUNE	BINDING SITE; CHAIN: B,	PROMYELOCYTIC	LEUKEMIA ZINC FINGER PROTEIN PLZF: CHAIN: A:				•					DNA; CHAIN: A, B, D, E;	PROTEIN; CHAIN: C, F, G;				TFILLA: CHAIN: A. D. 5S	RIBOSOMAL RNA GENE;	CHAIN: B, C, E, F;					YY1; CHAIN: C; ADENO-	ASSOCIATED VIKUS PS	DNA; CHAIN: A, B;
SEQ FOLD score		81.72			77.84										9.5					79.59	\ \ \						86.70		
PMF score				•		- ·																	٠			٠			
Verify						-			•															-					
Psi Blast		1e-29			8.5e-19			,			•	,			3.4e-48					5.16-30							1.76-32		
END		613			143										612				******	609	}						612		
START AA		531			19	-			-			•			230					437	· •						504		
CHAIN		A	-		Ą		,								ນ	•				V	:						C		
BOY ED		laih			Ibuo										lmey					1#6	2						lubd		
SEQ ID NO:		143			143						•				143	•				143	}						143		

PDB annotation	ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA- PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE- FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	SYNTHASE DHDPS; SYNTHASE, DIHYDRODIPICOLINATE	LYASE	KINASE KINASE, SIGNAL TRANSDUCTION, CALCIUM/CALMODULIN	TRANSFERASE TRANSFERASE, SERINE/THREONINE- PROTEIN KINASE, CASEIN KINASE, 2 SER/THR KINASE	the bank limb lime of land who lime there limes
Compound		ZINC FINGER PROTEIN GLI1; CHAIN: A; DNA; CHAIN: C, D;	DIHYDRODIPICOLINATE SYNTHASE; CHAIN: A, B;	N-ACETYLNEURAMINATE LYASE; INAL 4 CHAIN: 1, 2, 3, 4; INAL 5	CALCEUM/CALMODULIN- DEPENDENT PROTEIN KINASE; CHAIN: NULL;	PROTEIN KINASE CK2/ALPHA-SUBUNIT; CHAIN: NULL;	TRANSFERASE(PHOSPHO TRANSFERASE) \$C./AMP\$- DEPENDENT PROTEIN KINASE (E.C.2.7.1.37) (\$C/APK\$) 1APM 3 (CATALYTIC SUBUNIT) ALPHA ISOENZYME MUTANT WITH SER 139 1APM 4 REPLACED BY ALA (\$1394\$) COMPLEX WITH THE PEPTIDE 1APM 5 INHIBITOR PKI(5-24) AND THE DETERGENT
SEQ FOLD score		71.69	113.45	97.53	29.86	129.78	115.15
PMF							
Verify		·		·			
Psi Blast		1.7e-31	3.4e-83	1.2e-55	8.5e-60	1e-86	8.5e-48
END		[19	272	268	510	83	533
START AA		473			209	187	183
CHAIN		¥.	4	1			ш
PDB ID		2gli	1dhp	lnal	1a06	1a60	lapm
SEQ ID NO:		143	154	154	157	157	

																			······································		التسار كا		
PDB annotation		PROTEIN KINASE CDK2; PROTEIN KINASE, CELL	PHOSPHORYLATION, CTAIRDORDORNE 2 CELT	DIVISION, MITOSIS, INHIBITION	COMPLEX	(KINASE/INHIBITOR) CDK6; P19INK4D; CYCLIN	DEPENDENT KINASE,	KINASE INHIBITORY 2	PROTEIN, CDK, INK4, CELL	(KINASE/INHIBITOR) HEADER HELIX	COMPLEX (INHIBITOR PROTEIN/KINASE)	INHIBITOR PROTEIN,	KINASB, CELL CYCLE 2	CONTROL, ALPHA/BETA, TO COMPLEX (INHIBITOR	PROTEIN/KINASE)			· · · · · · · · · · · · · · · · · · ·			B. Um	Genz E.	
Compound	MEGA-8 1APM 6	CYCLIN-DEPENDENT PROTEIN KINASE 2;	CIPCIN: NOVE		CYCLIN-DEPENDENT	KINASE 6; CHAIN: A, C; CYCLIN-DEPENDENT	KINASE INHIBITOR; CHAIN: B. D:				CYCLIN-DEPENDENT KINASH 6: CHAIN: A:	P19INK4D; CHAIN: B;		:		PHOSPHOTRANSFERASE CAMP-DEPENDENT	PROTEIN KINASE CATALYTIC SUBUNIT	1CMK 3 (B.C.2.7.1.37)	TRANSFERASE (PHOSPHO	DEPENDENT PROTEIN	KINASE (E.C.2.7.1.37) (CAPK) 1CTP 3	(CATALYTIC SUBUNIT)	
SEQ FOLD score		160.34			148.04						163.82			•		116.16			112.97			•	
PMF score												-					,						
Verify score		·																-			<u>.</u>	:	
Psi Blast		0		:	1.2e-91						6.8e-		,			5.1e-49		·	6.8e-50				
A END		524			513	•			•		83					533			509			•	
START AA		214		!	214						210					163		•	181				
CHAIN			,	•	Ą						¥					m)			E				
POB BOB		laq1		٠.	1bi8						1blx			,		Icmk			lctp				
SEQ ID NO:		157			157						157		_			157			157				

Table 5

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PDB annotation	PROTEIN KINASE CDK2; TRANSFERASE,	SERINE/THREONINE	PROTEIN KINASE, ATP-	BINDING, 2 CELL CYCLE,	PHOSPHORYLATION	SERINE/THREONINE-	PROTEIN KINASE CSBP, RK,	P38; PROTEIN SER/THR-	KINASE,	SERINE/THREONINE.	TO ANGEGO A GE TAIK 3.	TRANSFERASE INK3 MAP	KINASE,	SERINE/THREONINE	PROTEIN 2 KINASE	KINASE KINASE, TWITCHIN,	INTRASTERIC REGULATION	KINASE KINASE, TWITCHIN,	INTRASTERIC REGULATION	TRANSFERASE MITOGEN	ACTIVATED PROTEIN	KINASE; TKANSFEKASE,	MAP KINASE,	PROTEIN KINASE 2 P38	KINASE RABBIT MUSCLE	PHOSPHORYLASE KINASE;	GLYCOGEN METABOLISM, 🖺	TRANSFERASE,	SERINE/THREONINE	PROTEIN, 2 KINASE, ATP-	BINDING, CALMODULIN-	TRANSFERASE MAP	KINASE, SERINE/THREONINE
Compound	HUMAN CYCLIN- DEPENDENT KINASE 2;	CHAIN: NULL;		,		P38 MAP KINASE: CHAIN:	NULL;				C. ITIN N. TEDMINAT	KINASE: CHAIN: NULL:	•			TWITCHIN; CHAIN: NULL;	•	TWITCHIN; CHAIN: A, B;	•	MAP KINASE P38; CHAIN:	NULL;		•		PHOSPHORYL ASE	KINASE; CHAIN: NULL;						ERK2; CHAIN: NULL;	
SEQ FOLD score	171.50					145.96					162.87		-			133.78		134.02		174.44					96.25							161.09	
PMF score					-								•																	,			•
Verify score				·.																													
Psi Blast	. 0				•	0					0	,				8.5e-57		8.5e-54		0					6.8e-57						. <u></u>	0	
	524					120					544	:	•			629		540		260					482							557.	
START AA	214					200					200	}				192		187		199					214							210	
CHAIN																		✓									•						
PDB ED	1hcl					lian				· · · · ·	1iik	ļ				1koa		1kob		1p38					1phk	•						1pme	·
SEQ ID NO:	157				•	157					157					157		157		157			-		157							157	

					<u> </u>	 The state of the second	- 17 17 17 most
PDB annotation	PROTEIN KINASE, TRANSFERASE	SERINE KINASE SERINE KINASE, TITIN, MUSCLE, AUTOINHIBITION	TRANSFERASE MITOGEN ACTIVATED PROTEIN KINASE, MAP 2, ERK2; TRANSFERASE, SERINE/THREONINE- PROTEIN KINASE, MAP KINASE, 2 ERK2	COMPLEX (NUCLEAR PROTEIN/RNA) COMPLEX (NUCLEAR PROTEIN/RNA), RNA, SNRNP,RIBONUCLEOPROTE IN	COMPLEX (NUCLEAR PROTEINIRNA) COMPLEX (NUCLEAR PROTEINIRNA), RNA, SNRNP, RIBONUCLEOPROTE "	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER, DNA PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER, DNA)
Compound		TITIN; CHAIN: A, B;	EXTRACELLULAR REGULATED KINASE 2; CHAIN: NULL;	U2 RNA HARRPIN IV; CHAIN: Q. R; U2 A; CHAIN: A, C; U2 B"; CHAIN: B, D;	U2 RNA HAIRPIN IV; CHAIN: Q, R; U2 A; CHAIN: A, C; U2 B"; CHAIN: B, D;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;
SEQ FOLD score		127.01	165.77	54.53	53.48	 74.63	97.41
PMF score						·	
Verify							
Psi Blast		1.46-45	o [·]	3.46-28	1.26-28	5.1e-30	1.4c-48
END		534	553	304	317	470	329
START AA		210	198	136	136	388	247
CHAIN		4		V	ပ	¥	ပ
PDB U		145	3erk	1a9n	la9n	lalh	1mey
SEQ ID NO:		157	157	163	163	168	168

						
PDB annotation	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING- YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA- PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE- FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	HYDROLASE TETRATRICOPEPTIDE, TRP; HYDROLASE, PHOSPHATASE, PROTEIN- PROTEIN INTERACTIONS, TPR, 2 SUPER-HELIX, X-RAY STRUCTURE	TRANSPORT PROTEIN RHO- GTPASE EXCHANGE FACTOR, TRANSPORT PROTEIN	PHOSPHOTRANSFERASE C- SRC, P60-SRC; SRC, TYROSINE KINASE, PHOSPHORYLATION, SH2,
Compound	TFIIIA; CHAIN: A, D; SS RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	SERINB/THREONINE PROTEIN PHOSPHATASE 5; CHAIN: NULL;	PIX; CHAIN: A;	TYROSINE-PROTEIN KINASE SRC; CHAIN: NULL;
SEQ FOLD score	104.01	91.10	99.34	96.15	99.42	73.30
PMF score			·			·
Verify						
Psi Blast	5.1e-37	3.4e-35	1.26-34	5.1e-22	1.2e-30	1.7e- 100
END	384	581	358	430	362	495
START	216	473	222	274	150	70
CHAIN	₹	U	¥		∀	
PDB ED	1466	1ubd	2gli	lai7	1by1	1fmk
SEQ ID NO:	168	168	168	171	173	173

Table 5

				 ,	_	##: ## ·	
PDB annotation	SH3, 2 PHOSPHOTYROSINE, PROTO-ONCOGENE, PHOSPHOTRANSFERASE	TYROSINE KINASE TYROSINE KINASE. INHIBITOR COMPLEX, DOWN-REGULATED KINASE, 2 ORDERED ACTIVATION LOOP	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	ENDONUCLEASE ENDONUCLEASE, TRNA ENDONUCLEASE		KINASE KINASE, SIGNAL TRANSDUCTION, CALCIUM/CALMODULIN	
Compound		HAEMATOPOETIC CELL KINASE (HCK); CHAIN: A;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	TRNA ENDONUCLEASE; CHAIN: A, B, C, D;		CALCIUM/CALMODULIN- DEPENDENT PROTEIN KINASE; CHAIN: NUIL;	TRANSFERASE(PHOSPHO TRANSFERASE) \$C-/AMP\$- DEPENDENT PROTEIN KINASE (B.C.2.7.1.37) (\$C/APK\$) 1APM 3 (CATALYTIC SUBUNIT) ALPHA ISOENZYME MUTANT WITH SER 139 1APM 4 REPLACED BY ALA (\$139A\$) COMPLEX WITH THE PEPTIDE 1APM 5 INHIBITOR PKI(5-24) AND THE DETERGENT MEGA-8 1APM 6
SEQ FOLD score		74.45	73.78	61.23		119.44	251.49
PMF score							
Verify							
Psi Blast		3.4e- 100	5.16-50	1.7e-42		3.4e-81	0
END	·	490		172		458	478
START AA		89	103	1		139	113
CHAIN	-	Ą	ပ	A			ជ
808 ED		lqcf	1теу	1a79		1a06	lapm
SEQ ID NO:		173	186	188		191	191

Table 5

					-													Harris II	-	بسو		ر		, b n	-	- 11	<u></u>	JE IE	IE	EI.
PDB annotation	PROTEIN KINASE CDK2; PROTEIN KINASE, CELL CVCT R	PHOSPHORYLATION,	STAUROSPORINE, 2 CELL DIVISION, MITOSIS,	INHIBITION	COMPLEX (KINASE/INHIBITOR) CDK6;	P19INK4D; CYCLIN	DEPENDENT KINASE,	CYCLIN DEPENDENT	PROTEIN, CDK, INK4, CELL	CYCLE, COMPLEX	(KINASE/INHIBITOR)	HEADER HELLX	COMPLEX (INHIBITOR	PROTEIN/KINASE) INHIBITOR PROTEIN	CYCLIN-DEPENDENT	KINASE, CELL CYCLE 2	CONTROL, ALPHA/BETA,	COMPLEX (INHIBITOR PROTEIN/KINASE)			•			-					7	PROTEIN KINASE CDK2;
Compound	CYCLIN-DEPENDENT PROTEIN KINASE 2; CHAIN: MIT I	CITAIN: INOLL,			CYCLIN-DEPENDENT KINASE 6: CHAIN: A. C.	CYCLIN-DEPENDENT	KINASE INHIBITOR;	CHAIN: B, D;	•				CYCLIN-DEPENDENT	KINASE 6; CHAIN: A;					PHOSPHOTRANSFERASE	CAMP-DEPENDENT	PROTEIN KINASE	CATALYTIC SUBUNIT	ICMK 3 (E.C.2.7.1.37)	TRANSFIRASF/PHOSPHO	TRANSFERASE) CAMP-	DEPENDENT PROTEIN	KINASE (E.C.2.7.1.37)	(CAPK) 1CTP 3	CALALI IIC SUBUNII)	HUMAN CYCLIN-
SEQ FOLD score	110.89		•		112.00								135.09						252.68					244.28						123.56
PIMF score											-			-	.•															
Verify score																										٠				
Psi Blast	5.1e-51				3.4e-39								1.5e-42						0					c	,					1.7e-53
END	432				458	•							465						478					450	}				•	432
START	145				146			•		•			140						105					110	2					145
CHAIN					Ą			,					¥			•			H				٠,	Д	}					
20 E	laq1	 -			1bi8							7	1blx	<u> </u>		_			1cmk				•	<u> </u>						1hcl
SEQ ID	161				191		•						191						161	_				101						191

						·		
PDB annotation	TRANSFERASE, SERINE/THREONINE PROTEIN KINASE, ATP- BINDING, 2 CELL CYCLE, CELL DIVISION, MITOSIS, PHOSPHORYLATION	KINASE KINASE, TWITCHIN, INTRASTERIC REGULATION	TRANSFERASE MITOGEN ACTIVATED PROTEIN KINASE: TRANSFERASE, MAP KINASE, SERINE/THREONINE- PROTEIN KINASE, 2 P38	KINASE RABBIT MUSCI.E PHOSPHORYLASE KINASE; GLYCOGEN METABOLISM, TRANSFERASE, SERINE/THREONINE- PROTEIN, 2 KINASE, ATP- BINDING, CALMODULIN- BINDING	SERINE KINASE SERINE KINASE, TITIN, MUSCI.E, A	TRANSFERASE MITOGEN ACTIVATED PROTEIN KINASE, MAP 2, ERK2; TRANSFERASE, SERINE/THREONINE- PROTEIN KINASE, MAP	OXYGEN TRANSPORT OXYGEN TRANSPORT	the Broke Heart Cont
Compound	DEPENDENT KINASE 2; CHAIN: NULL;	TWITCHIN; CHAIN: A, B;	MAP KINASE P38; CHAIN: NULL;	PHOSPHORYLASB KINASB; CHAIN: NULL;	TITIN; CHAIN: A, B;	EXTRACELLULAR REGULATED KINASE 2; CHAIN: NULL;	HEMOGLOBIN; CHAIN: A, E, C, F;	OXYGEN TRANSPORT HEMOGLOBIN (DEOXY, HUMAN FETAL F=/II\$=) IFDHG 1 IFDHH 2
SEQ FOLD score		134.48	119.00	132.88	114.63	121.20	76.31	67.75
PMF score								
Verify score		·						
Psi Blast		5.1e-64	3.4e-46	1.7e-74	3.4e-52	3.4e-43	6.8e-30	3.4e-29
END		488	508	424	475	206	78	78
START AA		118	128	144	142	132	2	£
CHAIN ID		A			¥		EX	9
PDB ID		1kob	1p38	1phk	14ki	3erk	1a9w	Ifdh
SEQ ID NO:		191	191	191	161	191	193	193

		δ. m. γ.	ZZ ~	~ * * *	ZZ	7.07	S
uo		COMPLEX (GTP- BINDING/TRANSDUCER) BETA1, TRANSDUCIN BETA SUBUNIT; GAMMA1, TRANSDUCIN GAMMA SUBUNIT; COMPLEX (GTP- BINDING/TRANSDUCER), G PROTEIN, HETEROTRIMER 2 SIGNAL TRANSDUCTION	COMPLEX (TRANSCRIPTION REGULATION/DNA) GABPALPHA, GABPBETAI; COMPLEX (TRANSCRIPTION REGULATION/DNA), DNA- BINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, ANKYRIN REPRATS, TRANSCRIPTION 3 PACTOR	TRANSCRIPTION FACTOR P65; P50D; TRANSCRIPTION FACTOR, IKB/NFKB COMPLEX	COMPLEX (TRANSCRIPTION REC/ANK REPEAT) COMPLEX (TRANSCRIPTION REGULATION/ANK REPEAT), ANKYRIN 2 REPEAT HELIX	OXIDOREDUCTASB TRYPANOTHIONB REDUCTASE, FAD DEPENDENT DISULPHIDE 2 OXIDOREDUCTASE	COMPLEX (OXIDOREDUCTASE/TRANS
PDB annotation		COMPLEX (GTP-BINDING/TRANSDUCER) BETA1, TRANSDUCIN BESUBUNT; GAMMA1, TRANSDUCIN GAMMA SUBUNT; COMPLEX (GTBINDING/TRANSDUCER) PROTEIN, HETEROTRIME	COMPLEX (TRANSCRIP REGULATION/DNA) GABPALPHA; GABPBET COMPLEX (TRANSCRIP REGULATION/DNA), DN REGULATION/DNA), DN BINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, ANKYRIN REPRATS, TRANSCRIPTION 3 PACT	TRANSCRIPTION FACTOR P65; P50D; TRANSCRIPTIO FACTOR, IKB/NFKB COMPLEX	COMPLEX (TRANSCR REG/ANK REPEAT) COMPLEX (TRANSCR REGULATION/ANK REPEAT), ANKYRIN 2 REPEAT HELIX	OXIDOREDUCTASE TRYPANOTHIONE REDUCTASE, FAD DEPENDENT DISUL OXIDOREDUCTASE	CTASI
PDB a	٠	COMPLEX (GTP- BINDING/TRANS BETA1, TRANSD SUBUNIT; GAM TRANSDUCIN G, SUBUNIT; COMP BINDING/TRANS PROTEIN, HETEI	LEX (T LATIO) ALPHA LEX (T LATIO) NG, 2 N SIN, ET SCRIPT	TRANSCRIPTION FA P65; P50D; TRANSCI FACTOR, IKB/NFKB COMPLEX	COMPLEX (TRANS) REG/ANK REPEAT) COMPLEX (TRANS) REGULATION/ANK REPEAT), ANKYRII REPEAT HELIX	OXIDOREDUCTAS TRYPANOTHIONE REDUCTASE, FAD DEPENDENT DISU OXIDOREDUCTAS	CEX
		COMP BETA: BETA: SUBUJ TRAN; SUBUJ BINDI PROTI	COMP CABP, CABP, COMP REGUI BINDI PROTI	TRANSCRI P65; P50D; FACTOR, I COMPLEX	COMP COMP REGU REPEA	OXIDO TRYPA REDUA DEPEN	COMPLEX (OXIDORE
		GT.	N N ETA AIN:	FE .		A, B;	
puno		ALPHA AIN: A; B; GT- IN: G;	ROTEII F. A; GA TEIN BI NA; CH	65 JN: A; SUBUJ PPA-B	65; CH/ A-B P50 KAPPA I: B, F;	NE HAIN:	AMIDE IASE;
Compound		HA/GI- VA; CHA HAIN; Y; CHA)	DING P CHAIN G PROT N: B; D	PA-B P II; CHA B P50D C; I-KA CHAIN	PA-B P -KAPP, B, D; I- CHAIN	VOTHIC	ROGEN
		GT-ALPHA/GI-ALPHA CHIMERA; CHAIN: A; GT- BETA; CHAIN: B; GT- GAMMA; CHAIN: G;	GA BINDING PROTEIN ALPHA; CHAIN: A; GA BINDING PROTEIN BETA 1; CHAIN: B; DNA; CHAIN: D, E;	NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF- KAPPA-B P50D SUBUNIT; CHAIN: C; I-KAPPA-B- ALPHA; CHAIN: D;	NP-KAPPA-B P65; CHAIN: A, C; NF-KAPPA-B P50; CHAIN: B, D; I-KAPPA-B- ALPHA; CHAIN: B, F;	TRYPANOTHIONE REDUCTASE; CHAIN: A, B;	DIHYDROLIPOAMIDE DEHYDROGENASE;
OLD re	٠						
SEQ FOLD score		66.17	94.15	83.26	91.60	78.99	115.94
PMF score					·		
Verify score							
Psi Blast		1.5e-45	1.4e-39	3.4e-40	6.8e-41	1e-58	1.5e-91
END		311	169	262	249	597	597
START AA		I				149	157
CHAIN							
	_	<u> </u>	M	Δ	<u>m</u>	₹	∢ .
PDB ID		1got	lawc	1.1km	Infi	laog	1ebd
SEQ ID NO:		195	199	661	199	71	17
J 2		-	H .	21	=	217	217

Table

PDB annotation	FERASE) E3BD; REDOX- ACTIVE CENTER, GLYCOLYSIS, OXIDOREDUCTASE		OXIDOREDUCTASE REDOX- ACTIVE CENTER, OXIDOREDUCTASE, FLAVOPROTEIN, FAD, NADP	the line of the line there the line is the line there there the line is the line there there is the line is the li	£
Compound	CHAIN: A, B; DIHYDROLIPOAMIDB ACETYLTRANSFERASE; CHAIN: C;	ELECTRON TRANSPORT (FLAVOCYTO CHROME) FLAVOCYTOCHROME C SULFIDE DEHYDROGENASE (FCSD) 1FCD 3	TRYPANOTHIONE REDUCTASE; CHAIN: A, B;	OXIDOREDUCTASE(FLAV OENZYME) GLUTATHIONE REDUCTASE (E.C.1.6.4.2) NAD MUTANT WITH ALA 179 IGES 3 REPLACED BY GLY, ALA 183 BY GLY, VAL 197 BY GLY, VAL 197 BY GLY, ALA 183 BY GLY, VAL 197 BY GLY, VAL 197 BY GLY, ALA 183 BY GLY, VAL 197 BY GLY, VAL 197 BY GLY, VAL 197 BY GLY, VAL 199 BY HEE 1GES 5 (A179G-A183G,V197E,R198 M,K199F,HZ00D,R204P) COMPLEXED WITH 1GES 6 NAD 1GES 7 OXIDOREDUCTASE DIHYDROGENASE GEC.1.8.1.4) COMPLEX WITH 1LPF 3 FLAVIN- ADENINE-DINUCLEOTIDE (FAD) 1LPF 4 OXIDOREDUCTASE	
SEQ FOLD score		77.29	91.61	100.73	
PMIF score	•				
Verify score					
Psi Blast		0.00051	1.7e-58	1.4e-77 1.4e-77 3.4e-83 5.1e-70	
END		<i>578</i>	597	593	
START AA		193	132	160	
CHAIN		A	V	4	
PDB ID		1fcd	1fec	lges lipf	
SEQ ID NO:		217	217	217 217	

				<u> </u>	The state of the s	all with their their their
PDB annotation			OXIDOREDUCTASE REDOX- ACTIVE CENTER, GLYCOLYSIS, OXIDOREDUCTASE, NAD, 2 FLAVOPROTEIN, FAD, P64K		OXIDOREDUCTASE NTR; HYPOTHETICAL PROTEIN, REDOX-ACTIVE CENTER, OXIDOREDUCTASE, 2 DISULTIDE OXIDOREDUCTASE, THIOREDOXIN REDUCTASE, FLAVIN 3 ADENINE DINULEOTIDE	
Compound	DIHYDROLIPOAMIDE DEHYDROGENASE (E.C.1.8.1.4) COMPLEX WITH 1LVL 3 NICOTINAMIDE- ADENINE-DINUCLEOTIDE (NAD+) 1LVL 4	OXIDOREDUCTASE (#202(A)) NADH PEROXIDASB (NPX) (E.C.1.11.1.1) MUTANT WITH CYS 42 INHP 3 REFLACED BY ALA (C42A) INHP 4	SURFACE PROTEIN; CHAIN: NULL;	OXIDOREDUCTASE(FLAV OENZYME) THIOREDOXIN REDUCTASE (E.C.1.64.5) MUTANT WITH CYS 138 1TRB 3 REFLACED BY SER (C138S) 1TRB 4	NADPH DEPENDENT THIOREDOXIN REDUCTASE; CHAIN: NULL;	OXIDOREDUCTASE (FLAVOENZYME) GLUTATHIONE REDUCTASE (B.C.1.6.4.2), OXIDIZED FORM (E) 3GRS
SEQ FOLD score		157.16	81.98	91.34	92.67	97.61
PMF score	•					
Verify						
Psi Blast		3.4e-64	5.1e-76	1.26-40	5.1e-39	8.5e-71
END		297	597	490	495.	593
START		195	147	190	174	158
CHAIN	-					
PDB CD		Inhp	lojt	<u> </u>	1vdc	3grs
SEQ ID		217	217	217	217	217

rable :

			 										 		-64-			1.1	liber c	i u	JE (_ []
PDB annotation			KINASE KINASE, SIGNAL TRANSDUCTION, CALCIUM/CALMODULÍN										COMPLEX	(KINASE/INHIBITOR) CDK6; P19INK4D; CYCLIN	DEPENDENT KINASE,	KTNASE INHIBITORY 2	PROTEIN, CDK, INK4, CELL	CYCLE, COMPLEX	(KINASE/INHIBITOR) HEADER HELIX				
Compound	4	OXIDOREDUCTASE DIHYDROLIPOAMIDE DEHYDROGENASE (E,C.1.8.1.4) 3LAD 3	CALCIUM/CALMODULIN- DEPENDENT PROTEIN KINASE; CHAIN: NULL;	TRANSFERASE(PHOSPHO TRANSFERASE) \$C-/AMP\$-	KINASE (E.C.2.7.1.37)	(\$C/APK\$) 1APM 3	(CATALYTIC SUBUNIT) ALPHA ISOENZYME	MUTANT WITH SER 139	1APM 4 REPLACED BY	WITH THE PEPTIDE 1APM	5 INHIBITOR PKI(5-24)	AND THE DETERGENT MEGA-8 1APM 6	CYCLIN-DEPENDENT	KINASE 6; CHAIN: A, C; CYCLIN-DEPENDENT	KINASE INHIBITOR;	Circuit: D, D,				PHOSPHOTRANSFERASE CAMP, DEPENDENT	PROTEIN KINASE	CATALYTIC SUBUNIT	ICMK 4
SEQ FOLD score		114.03	106.55	104.03									72.81						•	100.79			
PMF																							
Verify					•	•												•					
Psi Blast		3.4e-82	8.5e-76	6.8e-95	•					•		•	3.4e-38							8.5e-96			
END AA		597	245	254									235							254			
START AA		151	1	11-4									1							1			
CHAIN		Ą	•	я									A					-		В		_	
PDB ID		3lad	1a06	lapm _.									1bi8				,			1cmk			
SEQ ID NO:		217	219	219									219							219			

		•			•		•
PDB annotation		PROTEIN KINASE CDK2; TRANSFERASE, SERINE/THREONINE PROTEIN KINASE, ATP- BINDING, 2 CELL CYCLE, CELL DIVISION, MITOSIS, PHOSPHORYLATION	KINASE KINASE, TWITCHIN, INTRASTERIC REGULATION	KINASE RABBIT MUSCLE PHOSPHORYLASE KINASE; GLYCOGEN METABOLISM, TRANSFERASE, SERINE/THREONINE- PROTEIN, 2 KINASE, ATP- BINDING, CALMODULIN- BINDING	TRANSFERASE MAP KINASE, SERINE/THREONINE PROTEIN KINASE, TRANSFERASE	SERINE KINASE SERINE KINASE, TITIN, MUŞCLE, Ç AUTOINHIBITION	TRANSMEMBRANE PROTEIN COLICIN, BACTERIOCIN, ION CHANNEL FORMATION, TRANSMEMBRANE 2
Compound	TRANSFERASE(PHOSPHO TRANSFERASE) CAMP- DEPENDENT PROTEIN KINASE (E.C.2.7.1.37) (CAPK) 1CTP 3 (CATALYTIC SUBUNIT) 1CTP 4	HUMAN CYCLIN. DEPENDENT KINASE 2; CHAIN: NULL;	TWITCHIN; CHAIN: A, B;	PHOSPHORYLASB KINASB; CHAIN: NULL;	ERK2; CHAIN: NUIL;	TITIN; CHAIN: A, B;	COLICIN IA; CHAIN: NULL;
SEQ FOLD score	100.24	78.81	104.78	110.63	73.28	75.92	158.52
PMF score							
Verify							
Psi Blast	5.1e-91	3.4e-48	3.4e-59	1.4e-73	16-41	6.8e-48	6.8e-05
END AA	250	247	280	208	270	255	755
START AA	-	-	1	1	1		167
CHAIN	ш		Ą			А	
20g 80 ea	Ісф	Ihcl	1kob	1рћк	1pme	1175	1cii
SEQ ID NO:	219	219	219	219	219	219	222

Table 5

PDB annotation		HEPARIN-BINDING GROWTH FACTOR HEPARIN-BINDING GROWTH FACTOR		KINASE RI(ALPHA); REGULATORY SUBUNIT, KINASE		TRANSFERASE PIPK; LIPID SIGNALING, TRANSFERASE	CELL ADHESION PROTEIN EPITHELIAL CADHERIN DOMAINS 1 AND 2, ECAD12; CADHERIN, CELL ADHESION PROTEIN, CALCIUM BINDING PROTEIN	CELL ADHESION PROTEIN TO CELL ADHESION PROTEIN	Lags	INSECT IMMUNITY INSECT - IMMUNITY, LPS-BINDING, '- HOMOPHILIC ADHESION		CELL ADHESION PROTEIN
Compound		MIDKINE; CHAIN: A;	•	CAMP DEPENDENT PROTEIN KINASB; CHAIN: NUIL;		PHOSPHATIDYLINOSITOL PHOSPHATE KINASE IBETA; CHAIN: A, B;	E-CADHERIN; CHAIN: A, B;	N-CADHERIN; CHAIN: A;		HEMOLIN; CHAIN: A, B;	NEURAL ADHESION MOLECULE DROSOPHILA NEUROGLIAN (CHYMOTRYPTIC FRAGMENT CONTAINING THE ICFB 3 TWO AMINO PROXIMAL FIBRONECTIN TYPE II REPEATS 1CFB 4 (RESIDUES 610 - 814)) 1CFB 5	FIBRONECTIN; 1FNF 6
SEQ FOLD score		68.94		84.25		354.11	101.31	106.57		80.44	73.85	136.59
PMF score					٠	·						
Verify score										·		
Psi Blast		8.5e-22		1.5e-50		0	1.5e-20	6.8e-25		5.1e-27	8.5e-12	8.5e-42
END		139		724		421	1804	1811		398	345	435
START		76		454		39	1602	1609		1	166	25
CHAIN		A				A	¥	A		Ą		
PDB		1mkn		lrgs		1bo1	ledh	Incj		1bih	1cfb	1fnf
SEQ ID NO:		231		236.		241	242	242		244	244	244

Table 5

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PDB annotation	RGD, EXTRACELLULAR MATRIX 1FNF 18	HEPARIN AND INTEGRIN	BINDING HEPARIN AND INTEGRIN BINDING	STRUCTURAL PROTEIN	HEMIDESMOSOME,	FIBRONECTIN,	2 PROTEIN		UBIQUITIN CONTUGATION	CONTUGATION, UBIQUITIN-	CONTUGATING ENZYME	LIGASE E6AP; UBCH7;	BILOBAL STRUCTURE,	TRIOTETNITO 19	UBIQUITIN CONJUGATING	ENZYME	LIGASE UBIQUITIN,	BIOUTIN-CONJUGATING PROPERTY OF THE PROPERTY O	UBIOUTIN-CONJUGATING 4	ENZYME UBIQUITIN-	CONTUGATING ENZYME;	UBIQUITIN-CONJUGATING	ENZYME, UBIQUITIN-	DIRECTED 2 PROTEOLYSIS,	CELL CYCLE CONTROL,	UBIOUTTIN CONJUGATION	UBCI; UBIQUITIN	CONTUGATION, LIGASE	UBIQUITIN CONJUGATION UBIQUITIN CONJUGATION,
Compound	CHAIN: NULL; 1FNF.7	FIBRONECTIN; CHAIN: A;		INTEGRIN BETA-4	SUBUNII; CHAIN: A, B;	-			UBIQUITIN-	RAD6; CHAIN: A, B, C;		UBIQUITIN-PROTEIN	C. TELOTIFIER	C. OBIQUILIN	E2; CHAIN: D;		UBIQUITIN	CONJUGATING ENZYME;	UBC9; CHAIN: NULL;							UBIOUITIN	CONJUGATING ENZYME;	CHAIN: NULL;	UBIQUITIN CONTUGATING ENZYME;
SEQ FOLD score		83.24		91.86		٠	•		94.52			71.13					93.22		83.68					_		96.93			84.68
PMF score							,								•		•												
Verify score																													
Psi Blast	ŕ	5.1e-30		3.4e-20					5.1e-33			1.2e-24	•				3.4e-36		3.4e-28		-					6.8e-35			6.8e-30
END		435		247					258			251			·		253		252							250			256
START AA		163		57					26			103					66		95							98		1	1 1 1
CHAIN		A		∢					4		-	Ω					∢		A										
PDB ID		1fnh		1983					layz			lc4z					Іфоф		1u9a							2aak			2e2c
SEQ ID NO:		244		244					245			245		٠.			245		245							245			245

					•								
PDB annotation	UBIQUITIN CARRIER PROTEIN, THIOESTER 2 BOND, LIGASE	UBIQUITIN CONTUGATION UBC7; UBIQUITIN CONTUGATION, LIGASE, YEAST						÷				MICROTUBULES MICROTUBULES, ALPHA- TUBULIN, BETA-TUBULIN, GTPASE HELIX	ACTIN-BINDING PROTEIN, ACTIN-BINDING PROTEIN, CALCIUM-BINDING,
Compound	CHAIN: NULL;	UBIQUITIN CONIUGATING ENZYME; CHAIN: NULL;		LIPID-BINDING PROTEIN ADIPOCYTE LIPID-	BINDING PROTEIN COMPLEXED WITH ARACHIDONIC 1ADL 3 ACID 1ADL 4	LIPID-BINDING PROTEIN	PROTEIN (HUMAN	COMPLEXED IHMR 3 WITH ONE MOLECULE OF HIADITA ACTOR IHMR 4	RETINOL TRANSPORT CELLULAR RETINOL BINDING PROTEIN II (APO	FORM) (APO-CRBPII) 10PA	CELULAR LIPOPHILIC TRANSPORT PROTEIN P2 MYELIN PROTEIN (P2) IPMP 3	TUBULIN; CHAIN: A, B;	T-FIMBRIN; CHAIN: NULL;
SEQ FOLD score		88.78		83.63		90.26	÷		198.37		76.11	321.45	
PMF score													0.95
Verify score	·		1,								•		0.41
Psi Blast		1.2e-29		8.5e-54		5.1e-56			8.5e-49		3.4e-53	0	1.2e-20
END		250		294		294	•		294		294	281	179
START AA		97		162		162			162		162	1	22
CHAIN									Ą		¥	Ø	
808 E		2ucz		ladi		1hmr		٠.	lopa	-	lpmp	1mb	1aoa
SEQ ID NO:		245		247		247			247		247	249	255

Table

						_	Z.													11-11			ىت	-		- IV	1	TF	- 11.	II "	السنيسا		
PDB annotation	PHOSPHORYLATION	ACTIN-BINDING PROTEIN	CALCIUM-BINDING,	PHOSPHORYL ATION	ACTIN-BINDING CALPONIN HOMOLOGY (CH) DOMAIN:	FILAMENTOUS ACTIN-	BINDING DOMAIN,	CYTOSKELETON	STRUCTURAL PROTEIN	DYSTROPHIN, MUSCULAR	DYSTROPHY, CALPONIN	ACTIN-BINDING, UTROPHIN	STRUCTURAL PROTBIN	CALPONIN HOMOLOGY	DOMAIN, DOMAIN	SWAPPING, ACTIN	BINDING, 2 UTROPHIN,	DISIROPHIN, CTRICTION AT DEOTERN	STRUCTORAL PROTEIN	EXTRACELLULAR MODULE #	OSTEONECTIN, SPARC,	SECRETED PROTEIN ACIDIC	AND EXTRACELLULAR	MODULE, GLYCOPROTEIN,	ANTI-ADHESIVE PROTEIN, 24	DIBECTED MITTAGENESIS	GI.YCOSYI.ATED 3 PROTEIN	MODRES	4	CHAPERONE HSP40;	CHAPERONE, HEAT SHOCK, F	MOLECULAR CHAPERONE	HDJ-1; MOLECULAR
Compound		T-FIMBRIN; CHAIN: NULL;			SPECTRIN BETA CHAIN; CHAIN: A:				DYSTROPHIN; CHAIN: A,	B, C, D;			UTROPHIN ACTIN	BINDING REGION; CHAIN:	A.B;					BASEMENT MEMBRANE	PROTEIN BM-40; CHAIN:	A, B;					•			DNAJ; CHAIN: NULL;		HUMAN HSP40; CHAIN:	NOLL;
SEQ FOLD score															,					60.57													
PMF score		0.04	•		0.07				20.0-				-0.05											-		•				96.0		1.00	
Verify score		-0.09			0.03				0.21				0.03																	0.49		0.65	
Psi Blast		8.5e-06	-		0.00068				3.4e-31				6.8e-30							3.4e-20										5.1e-27		3.4e-27	
END		133			137				179	•			179	_			 -			435		,	٠.							340		341	
START		6			27	,			8			•	24							220										275	٠.	912	
CHAIN					∢				A				A							A													
PDB		laoa			15kr				ldxx				lqag							Into										1bq0		1hdj	
SEQ ID NO:		255			255				255				255							260					•					262		262	7

Table 5

	٠.	•					10.16 Hands - 11.		
PDB annotation	CHAPERONE	TOXIN BINDING PROTEIN TWO DOMAINS: BETA PROPELLER AND ALPHA/BETA FOLD	TRANSCRIPTION INHIBITOR BETA-PROPELLER	TRANSCRIPTION INHIBITOR BETA-PROPELLER	TRANSCRIPTION INHIBITOR BETA-PROPELLER	TRANSCRIPTION INHIBITOR BETA-PROPELLER	COMPLEX (GTP-BINDING/TRANSDUCER) BETA1, TRANSDUCIN BETA SUBUNIT; GAMMA1, TRANSDUCIN GAMMA SUBUNIT; COMPLEX (GTP-SUBUNIT; COMPLEX (GTP-BINDING/TRANSDUCER), G PROTEIN, HETEROTRIMER 2, SIGNAL TRANSDUCTION	COMPLEX (GTP-BINDING/TRANSDUCER) BETA1, TRANSDUCIN BETA SUBUNIT; GAMMA1, TRANSDUCIN GAMMA SUBUNIT; COMPLEX (GTP-SUBUNIT; COMPLEX (GTP-BINDING/TRANSDUCER), G PROTEIN, HETEROTRIMER 2, SIGNAL TRANSDUCTION	COMPLEX (GTP-BINDING/TRANSDUCER) BETA1, TRANSDUCIN BETA
Compound		TOLB PROTEIN; CHAIN: A;	TRANSCRIPTIONAL REPRESSOR TUP1; CHAIN: A. B. C;	TRANSCRIPTIONAL REPRESSOR TUP1; CHAIN: A, B, C;	TRANSCRIPTIONAL REPRESSOR TUP1; CHAIN: A, B, C;	TRANSCRIPTIONAL REPRESSOR TUP1; CHAIN: A, B, C;	GT-ALPHA/GI-ALPHA CHIMERA; CHAIN: A; GT- BETA; CHAIN: B; GT- GAMMA; CHAIN: G;	GT-ALPHA/GI-ALPHA CHIMERA; CHAIN: A; GT- BETA; CHAIN: B; GT- GAMMA; CHAIN: G;	GT-ALPHA/GI-ALPHA CHIMERA; CHAIN: A; GT- BETA; CHAIN: B; GT-
SEQ FOLD score									
PMF score		0.00	0.23	1.00	-0.18	-0.18	0.76	-0.13	-0.19
Verify		0.09	0.03	0.02	0.04	-0.00	0.38	0.17	0.13
Psi Blast		1.7e-05	6.8e-63	1.4e-69	1.2e-61	1.4e-56	1e-78	6.8e-69	6.8e-48
END AA		202	433	316	. 789	760	314	6.29	761
START AA		18	104	10	401	498	1	398	498
CHAIN		Ą	Ą	A	A	Ą	e e	В	В
PDB		lcrz	lerj	lerj	lerj	1erj	1got	1got	lgot
SEQ ID NO:		263	263	263	263	263	263	263	263

Cable 5

PDB annotation	SUBUNIT; GAMMA1, TRANSDUCIN GAMMA SUBUNIT; COMPLEX (GTP- BINDING/TRANSDUCER), G PROTEIN, HETEROTRIMER 2 SIGNAL TRANSDUCTION	COMPLEX (GTP-BINDING/TRANSDUCER) BETA1, TRANSDUCIN BETA SUBUNIT; GAMMA1, TRANSDUCIN GAMMA SUBUNIT; COMPLEX (GTP-BINDING/TRANSDUCER), G PROTEIN, HETEROTRIMER 2 SIGNAL TRANSDUCTION	COMPLEX (GTP- BINDING/TRANSDUCER) BETAI, TRANSDUCIN BETA SUBUNIT; GAMMAI, TRANSDUCIN GAMMA SUBUNIT; COMPLEX (GTP- BINDING/TRANSDUCER), G PROTEIN, HETEROTRIMER 2 SIGNAL TRANSDUCTION	OXIDOREDUCTASE ENZYME, NITRITE REDUCTASE, OXIDOREDUCTASE, DENITRIFICATION, 2 ELECTRON TRANSPORT, PERIFLASMIC	RIBOSOME 50S RIBOSOMAL. PROTEIN L2P, HMAL2, HL4; 50S RIBOSOMAL PROTEIN L3P, HMAL3, HL1; 50S RIBOSOMAL PROTEIN L4E, HMAL4, HL6; 50S RIBOSOMAL PROTEIN L5P,
Compound	GAMMA; CHAIN: G,	GT-ALPHA/GI-ALPHA CHIMERA; CHAIN: A; GT- BETA; CHAIN: B; GT- GAMMA; CHAIN: G;	GT-ALPHA/GI-ALPHA CHIMERA; CHAIN: A; GT- BETA; CHAIN: B; GT- GAMMA; CHAIN: G;	CYTOCHROME CD1 NITRITE REDUCTASE; CHAIN: A, B;	23S RRNA; CHAIN: 0; 5S RRNA; CHAIN: 9; RIBOSOMAL PROTEIN L2; CHAIN: A; RIBOSOMAL PROTEIN L3; CHAIN: B; RIBOSOMAL PROTEIN L4; CHAIN: C; RIBOSOMAL;
SEQ FOLD score			·		·
PMF score		0.19	-0.05	-0.17	0.94
Verify		0.28	0.10	0.13	-0.28
Psi Blast		5.1e-64	3.46-57	1.2e-22	3.6e-34
END A		355	430	221	98
START		51	8	4	81
CHAIN		M.	α	V	Ω
PDB		Igot	1got	1qks	1供
SEQ ID		263	263	263	266

Fable

PDB annotation	HMALS, HL13; 30S RIBOSOMAL PROTEIN HS6;	50S RIBOSOMAL PROTEIN	LI3P, HMALI3; 50S	KIBOSOMAL FROIBIN LI4F, HMAL14, HL27; 50S	RIBOSOMAL PROTEIN L15P,	HMAL15, HL9; 50S	RIBOSOMAL PROTEIN L18P,	HMAL18, HL12; 50S	HI 29, L19: 50S RIBOSOMAL	PROTEIN L19E, HMAL19,	HL24; 50S RIBOSOMAL	PROTEIN L21E, HL31; 50S	RIBOSOMAL PROTEIN L22P,	HMAL22, HL23; 50S	RIBOSOMAL PROTEIN L23P,	HMAL23, HL25, L21; 50S	RIBOSOMAL PROTEIN L24P,	HMAL24, HL16, HL15; 50S	RIBOSOMAL PROTEIN L24E,	HL21/HL22; 50S RIBOSOMAL	PROTEIN L29P, HMAL29,	HL33; 50S RIBOSOMAL	PROTEIN L30P, HMAL30,	HL20, HL16; 505	KUBUSUMAL FRUIEIN LSIE, 1	PROTEIN I 30E HI 5: 50S	RIBOSOMAL PROTEIN L37E	L35E: 50S RIBOSOMAL	PROTEINS L39E, HI 39E.	HL 46E; 50S RIBOSOMAL	PROTEIN LAE, LA, HLA; 50S	RIBOSOMAL PROTEIN LGP, 💾	HMAL6, HL10 RIBOSOME [1]	ASSEMBLY, RNA-RNA,	PROTEIN-RNA, PROTEIN-
Compound	PROTEIN LS; CHAIN: D; H		OTEIN	RIBOSOMAL PROTEIN H		ROTEIN		RIBOSOMAL PROTEIN H	OTEIN		ROTEIN		OTEIN	_	COTEIN	Ξ.	OTEIN		ROTEIN		ROTEIN		ROTEIN	LZ4; CHAIN: Q;		NIETO		ROTEIN		ROTEIN		OTEIN		NEL	L37AE; CHAIN: W; P
SEQ FOLD score																																			
PMG		• •		٠							•								•				٠				•								
Verify score																																			
Psi Blast																		****											-						
END AA											<u>.</u>																								
START AA																																			
CHAIN										_					•																				
PDB U							· ·																										_		
SEQ ID NO:																										-									

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PDB annotation	PROTEIN	RIBOSOME 50S RIBOSOMAL PROTEIN 1.2P, HMA12, HL4; 50S RIBOSOMAL PROTEIN 1.3P, HMA13, HL1; 50S RIBOSOMAL PROTEIN 1.4B, HMA14, HL6; 50S RIBOSOMAL PROTEIN 1.5P, HMAL5, HL13; 30S RIBOSOMAL PROTEIN 1.4P, HMAL14, HL27; 50S RIBOSOMAL PROTEIN 1.1SP, HMAL15, HL9; 50S RIBOSOMAL PROTEIN 1.1SP, HMAL18, HL12; 50S RIBOSOMAL PROTEIN 1.1SP, HMAL18, HL12; 50S RIBOSOMAL PROTEIN 1.1SP, HMAL18, HL12; 50S RIBOSOMAL PROTEIN 1.1SP, HMAL18, HL12; 50S RIBOSOMAL PROTEIN 1.1SP, HMAL18, HL12; 50S RIBOSOMAL PROTEIN 1.1SP, HMAL18, HL12; 50S RIBOSOMAL PROTEIN 1.1SP, HMAL18, HL12; 50S RIBOSOMAL PROTEIN 1.1SP, HMAL18, HL12; 50S RIBOSOMAL PROTEIN 1.1SP, HL29, L19; 50S RIBOSOMAL	PROTEIN L21E, HL31; 50S RIBOSOMAL PROTEIN L22P, HMAL22, HL23; 50S RIBOSOMAL PROTEIN L23P, HMAL23, HL25, L21; 50S RIBOSOMAL PROTEIN L24P, HMAL24, HL16, HL15; 50S RIBOSOMAL PROTEIN L24E, HMAL24, HL16, HL15; 50S RIBOSOMAL PROTEIN L24E, HL21/FIL22; 50S RIBOSOMAI
Compound	RIBOSOMAL PROTEIN L37E; CHAIN: X; RIBOSOMAL PROTEIN L39E; CHAIN: Y; RIBOSOMAL PROTEIN L44E; CHAIN: Z; RIBOSOMAL PROTEIN L6; CHAIN: 1	23S RRNA; CHAIN: 0; 5S RRNA; CHAIN: 9; RIBOSOMAL PROTEIN L2; CHAIN: A; RIBOSOMAL PROTEIN L3; CHAIN: B; RIBOSOMAL PROTEIN L4; CHAIN: C; RIBOSOMAL PROTEIN L5; CHAIN: D; RIBOSOMAL PROTEIN L7AE; CHAIN: E; RIBOSOMAL PROTEIN L10E; CHAIN: F; RIBOSOMAL PROTEIN L13; CHAIN: H; RIBOSOMAL PROTEIN L14; CHAIN: H; RIBOSOMAL PROTEIN L15; CHAIN: I; RIBOSOMAL PROTEIN L15; CHAIN: I; RIBOSOMAL PROTEIN L15; CHAIN: I; RIBOSOMAL PROTEIN L15; CHAIN: I; RIBOSOMAL PROTEIN L15; CHAIN: I; RIBOSOMAL PROTEIN	L18; CHAIN: K; RIBOSOMAL PROTEIN L18E; CHAIN: L; RIBOSOMAL PROTEIN L19; CHAIN: M; RIBOSOMAL PROTEIN L21E; CHAIN: N; RIBOSOMAL PROTEIN L21E; CHAIN: N; RIBOSOMAL PROTEIN L22; CHAIN: O; RIBOSOMAL PROTEIN
SEQ FOLD score			
PMF		0.70	
Verify		-0.22	
Psi Blast		8.56-21	
END		%	
START		∞	
CHAIN		Þ	
EQ EI			
SEQ ID NO:		266	

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PDB annotation	HL33; SOS RIBOSOMAL PROTEIN L30P, HMAL30, HL20, HL16; SOS RIBOSOMAL PROTEIN L31B, L34, HL30; SOS RIBOSOMAL PROTEIN L32B, HL5; SOS RIBOSOMAL PROTEIN L37B, L35E; SOS RIBOSOMAL PROTEINS L39E, HL39B, HL46E; SOS RIBOSOMAL PROTEIN L4B, LA, HL4; SOS RIBOSOMAL PROTEIN L6P, HMAL6, HL10 RIBOSOME ASSEMBLY, RNA-RNA, PROTEIN-RNA, PROTEIN- PROTEIN-RNA, PROTEIN- PROTEIN		SERINE PROTEASE PCPA2; SERINE PROTEASE, ZYMOGEN, HYDROLASE	SERINE PROTEASE PORCINE PROCARBOXYPEPTIDASE, SERINE PROTEASE	I	,	DNA BINDING PROTEIN CENTROMERE PROTEIN, DNA-BINDING, HELIX-
Compound	L23; CHAIN: P; RIBOSOMAL PROTEIN L24; CHAIN: Q; RIBOSOMAL PROTEIN L24E; CHAIN: R; RIBOSOMAL PROTEIN L29; CHAIN: S; RIBOSOMAL PROTEIN L30; CHAIN: T; RIBOSOMAL PROTEIN L31E; CHAIN: T; RIBOSOMAL PROTEIN L32E; CHAIN: Y; RIBOSOMAL PROTEIN L37E; CHAIN: Y; RIBOSOMAL PROTEIN L37E; CHAIN: X; RIBOSOMAL PROTEIN L37E; CHAIN: X; RIBOSOMAL PROTEIN L37E; CHAIN: X; RIBOSOMAL PROTEIN L39E; CHAIN: X; RIBOSOMAL PROTEIN L39E; CHAIN: X; RIBOSOMAL PROTEIN L39E; CHAIN: X; RIBOSOMAL PROTEIN L39E; CHAIN: X; RIBOSOMAL PROTEIN L39E; CHAIN: X; RIBOSOMAL PROTEIN L44E; CHAIN: Z; RIBOSOMAL PROTEIN L44E; CHAIN: Z; RIBOSOMAL PROTEIN L6; CHAIN: 1;		PROCARBOXYPEPTIDASB A2; CHAIN: NULL;	PROCARBOXYPEPTIDASE B; CHAIN: NULL	HYDROLASE(C- TERMINAL PEPTIDASE) PROCARBOXYPEPTIDASE A (B.C.3.4.12.2) 1PCA 3		CENTROMERE PROTEIN B; CHAIN: A;
SEQ FOLD score			107.01	110.63	114.78		
PMF score							0.83
Verify							0.08
Psi Blast			1.7e-91	1.7e-88	3.4e-90		5.1e-07
END			548	549	554		56
START			119	123	122		. 9
CHAIN							A
PDB			laye	Insa	Ipca		1bw6
SEQ ID			268	268	. 768		269

Table 5

							Antible prompt brown.
PDB annotation	TURN-HELIX, DNA 2 BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING FROTEIN		Constitution Cons
Compound		QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	TRANSCRIPTION REGULATION YEAST TRANSCRIPTION PACTOR ADRI (RESIDUES 102 - 130) 1ARD 3 (AMINO TERMINAL ZINC FINGER DOMAIN) (NMR, 10 STRUCTURES) 1ARD 4 (ADR1B) 1ARD 5	TRANSCRIPTION REGULATION YEAST TRANSCRIPTION FACTOR
SEQ FOLD score				·			
PMF score	·	0.27	0.81	0.11	0.25	0.00	0.3 <i>7</i>
Verify score		-0.37	0.00-	-0.17	-0.46	-0.13	-0.24
Psi Blast		8.5e-21	1.7e-28	1.7e-28	1.2e-17	7.2e-07	6.8e-05
END		301	412		466	278	280
START		224	333	361	361	252	252
CHAIN		А	¥	¥.	V		
PDB ID		1a1h	lalh	laih	lalh	lard .	lard
SEQ ID NO:		270	270	270	270	270	270

Sable 5

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PDB annotation				GENE REGULATION POZ DOMAIN; PROTEIN- PROTEIN INTERACTION DOMAIN, TRANSCRIPTIONAL 2 REPRESSOR, ZINC-FINGER PROTEIN, X-RAY CRYSTALLOGRAPHY, 3 PROTEIN STRUCTURE, PROMYELOCYTIC LEUKEMIA, GENE REGULATION	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)
Compound	ADRI (RESIDUES 102 - 130) 1ARD 3 (AMINO TERMINAL ZINC FINGER DOMAIN) (NMR, 10 STRUCTURES) 1ARD 4 (ADR1B) 1ARD 5	DNA-BINDING PROTEIN HUMAN ENHANCER- BINDING PROTEIN MBP-1 MUTANT WITH CYS 11 1BBO 3 REPLACED BY ABU (C11ABU) (NMR, 60 STRUCTURES) 1BBO 4	DNA-BINDING PROTEIN HUMAN ENHANCER- BINDING PROTEIN MBP-1 MUTANT WITH CYS 11 1BBO 3 REPLACED BY ABU (C11ABU) (NMR, 60 STRUCTURES) 1BBO 4	PROMYELOCYTIC LEUKEMÍA ZINC FINGER PROTEIN PLZP; CHAIN: A;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;
SEQ FOLD score					·
PMF score		0.40	0.58	0.71	0.33
Verify score		-0.22	-0.46	0.26	-0.31
Psi Blast		3.6e-07	3.6e-13	1.7e-37	3.4e-36
END		278	408	117	301
START AA		252	362	-	223
CHAIN				¥	ပ
PDB CD		1bbo	1bbo	1buo	1mey
SEQ TD NO:	·	270	. 270	270	270

Table 5

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PDB annotation	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERÂCTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURB, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZÎNC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	ZINC FINGER TRANSCRIPTION FACTOR SP1; ZINC FINGER, TRANSCRIPTION ACTIVATION, SP1	ZINC FINGER TRANSCRIPTION FACTOR SP1; ZINC FINGER, TRANSCRIPTION ACTIVATION, SP1	COMPLEX (TRANSCRIPTION
Compound	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, B; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, B; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	SPIF3; CHAIN: NULL;	SPIF2; CHAIN: NULL;	TRANSCRIPTION FACTOR
SEQ FOLD score							
PMF score	0.49	0.16	0.57	0.39	0.43	0.54	0.07
Verify	-0.13	-0.06	-0.45	-0.14	0.03	-0.40	0.15
Psi Blast	1.7e-48	8.5e-49	1.7e-12	3.4e-13	.3.6e-05	1.2e-07	5.1e-19
AA A	412	470	276	382	278	280	412
START AA	332	388	249	358 ·	252	252	333
CHAIN	ပ	ပ	ರ	ප		-	Ą
EGE CO	Imey	Imey	Imey	Imey	lspl	1sp2	11f3
SEQ ID NO:	270	270	270	270	270	270	270

Table 5

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PDB annotation	REGULATION/DNA) TFIIIA; 5S GENE; NMR, TFIIIA,	PROTEIN, DNA,	TRANSCRIPTION FACTOR,	SS KNA Z GENE, DNA BINDING PROTEIN. ZINC	FINGER, COMPLEX 3	TRANSCRIPTION	REGULATION/DNA)	COMPLEX (TRANSCRIPTION	KEGULATION/DINA) IFILIA;	PROTEIN DNA	TRANSCRIPTION FACTOR,	5S RNA 2 GENE, DNA	BINDING PROTEIN, ZINC	FINGER, COMPLEX 3	(TRANSCRIPTION	REGULATION/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA)	COMPLEX (TRANSCRIPTION	KEGULATION/DINA), KINA	POLYMERASE III, 2	IKANSCKIFIJON	PROTEIN	COMPLEX (TRANSCRIPTION	REGULATION/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA), RNA	POLYMERASE III, 2	TRANSCRIPTION	PROTRIN	COMPLEX (TRANSCRIPTION)	REGULATION/DNA) YING-	YANG 1; TRANSCRIPTION INTERIOR	
Compound	IIIA; CHAIN: A; 5S RNA GENE; CHAIN: E, F;							TRANSCRIPTION FACTOR	IIIA; CHAIN: A; 3S KNA	GEINE; CHAIN: E, F;							TFIIIA; CHAIN: A, D; 5S	RIBOSOMAL RNA GENE;	CHAIN: B, C, E, F;					TFIIIA; CHAIN: A, D; 5S	RIBOSOMAL RNA GENE;	CHAIN: B, C, E, F;					YY1; CHAIN: C; ADENO-	ASSOCIATED VIRUS P5	INITIATOR ELEMENT	M. M. Canada V. 1 24 20 1
SEQ FOLD score							٠				<u> </u>																							
PMF score								0.25									0.03							0.33							0.29			
Verify								09.0-									-0.50							-0.35							-0.28			
Psi Blast								2.4e-12			•						3.4e-37				<u></u>			1.7e-36							3.4e-34			
END AA	·				•			466									479			•			•	507							442			
START AA								361									333							361							340			
CHAIN								٧									¥	,						٧						٠.	ပ			
202 133 133 133 133 133 133 133 133 133 13								1163									1tf6							1466							Iubd			
SEQ ID NO:								270					•				270							270							270			T

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PDB annotation	ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA- PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	TRANSCRIPTION REGULATION TRANSCRIPTION REGULATION, ADRI, ZINC FINGER, NMR	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE- FINGER GLI, GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE- FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	SCAFFOLD PROTEIN, PP2A, PHOSPHORYLATION, HEAT REPEAT	TRANSPORT PROTEIN SERINE-RICH RNA POLYMERASE I SUPPRESSOR PROTEIN; ARM REPEAT	NUCLEAR IMPORT RECEPTOR KARYOPHERIN ALPHA; NUCLEAR IMPORT RECEPTOR, NUCLEAR LOCALIZATION SIGNAL, 2 ARMADILLO REPEATS, AUTOINHIBITION, INTRASTERIC REGULATION	SMALL GTPASE
Compound		ADR1; CHAIN: NULL;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLI1; CHAIN: A; DNA; CHAIN: C, D;	PROTEIN PHOSPHATASE PP2A; CHAIN: A, B;	KARYOPHERIN ALPHA; CHAIN: A, B; MYC PROTO- ONCOGENE PROTEIN; CHAIN: C, D, E, F;	IMPORTIN ALPHA; CHAIN: A;	RAN; CHAIN: A, C;;
SEQ FOLD score							·	
PMF score		0.12	0.09	0.11	0.21	0.24	0.25	0.07
Verify score		-0.69	-0.15	-0.53	-0.30	0.16	0.03	90.0
Psi Blast		3.4e-11	1.7e-07	1.2e-15	1.2e-27	1.4c-47	8.56-50	3.4e-20
AA AA		278	275	466	888	838	838	574
START AA	·	231	224	361	478	436	437	110
CHAIN			∀	A	А	⋖	< <	В
PDB CI		2adr	2gli	2gli	163u	1664	lial	libr
SEQ ID NO:		270	270	270	272	272	272	272

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PDB annotation	KARYOPHERIN BETA, P95 SMALL GTPASE, NUCLEAR TRANSPORT RECEPTOR	NUCLEAR TRANSPORT PROTEIN COMPLEX HEAT REPEATS, NUCLEAR TRANSPORT PROTEIN COMPLEX	TRANSPORT RECEPTOR KARYOPHERIN BETA-1, NUCLEAR FACTOR P97, IMPORTIN IMPORTIN ALPHA-2 SUBUNIT, KARYOPHERIN ALPHA-2 TRANSPORT RECEPTOR, NUCLEAR IMPORT, HEAT MOTIF, NLS-BINDING	TRANSPORT RECEPTOR KARYOPHERIN BETA-1, NUCLEAR FACTOR P97, IMPORTIN IMPORTIN ALPHA-2 SUBUNIT, KARYOPHERIN ALPHA-2 TRANSPORT RECEPTOR, NUCLEAR IMPORT, HEAT MOTIF, NIS-BINDING	STRUCTURAL PROTEIN ARMADILLO REPEAT, BETA-CATENIN, STRUCTURAL PROTEIN	SCAFFOLD PROTEIN SCAFFOLD PROTEIN, PP2A, PHOSPHORYLATION, HEAT REPEAT	SCAFFOLD PROTEIN, PP2A, PHOSPHORYLATION, HEAT
Compound	IMPORTIN BETA SUBUNIT; CHAIN: B, D	KARYOPHERIN BBTA2; CHAIN: B; RAN; CHAIN: C;	IMPORTIN BETA SUBUNIT; CHAIN: A; IMPORTIN ALPHA-2 SUBUNIT; CHAIN: B;	IMPORTIN BETA SUBUNIT; CHAIN: A; IMPORTIN ALPHA-2 SUBUNIT; CHAIN: B;	BETA-CATENIN; CHAIN: NULL;	PROTEIN PHOSPHATASE PP2A; CHAIN: A, B;	PROTEIN PHOSPHATASE PP2A; CHAIN: A, B;
SEQ FOLD score		116.75		127.17		125.57	
PMF			0.75		0.07		0.69
Verify score		·	-0.12		-0.06		-0.02
Psi Blast		1.7e-57	1.46-39	1.4e-39	2.46-17	1.2e-72	1.2e-72
END		828	894	849	709	850	847
START		က	109	0	239	961	207
CHAIN		Ø	V	V		4	V
PDB ID		1qbk	19gr	19gr	2bct	163u	1b3u
SEQ ID NO:		272	272	272	272	273	273

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PDB annotation	REPEAT	SCAFFOLD PROTEIN SCAFFOLD PROTEIN, PP2A, PHOSPHORYLATION, HEAT REPEAT	TRANSPORT PROTEIN SERINE-RICH RNA POLYMERASE I SUPPRESSOR PROTEIN; ARM REPRAT	NUCLEAR IMPORT RECEPTOR KARYOPHERIN ALPHA; NUCLEAR IMPORT RECEPTOR, NUCLEAR LOCALIZATION SIGNAL, 2 ARMADILLO REPEATS, AUTOINHIBITION, INTRASTERIC REGULATION	NUCLEAR TRANSPORT PROTEIN COMPLEX HEAT REPEATS, NUCLEAR TRANSPORT PROTEIN COMPLEX	TRANSPORT RECEPTOR KARYOPHERIN BETA-1, NUCLEAR FACTOR P97, IMPORTIN TMPORTIN ALPHA-2 SUBUNIT, KARYOPHERIN ALPHA-2 TRANSPORT RECEPTOR, NUCLEAR IMPORT, HEAT MOTIF, NLS-BINDING	HYDROLASE TETRATRICOPEPTIDE, TRP; HYDROLASE, PHOSPHATASE, PROTEIN-PROTEIN INTERACTIONS, TPP, 2 SUPER-HELIX, X-RAY
Compound		PROTEIN PHOSPHATASE PP2A; CHAIN: A, B;	KARYOPHERIN ALPHA; CHAIN: A, B; MYC PROTO- ONCOGENE PROTEIN; CHAIN: C, D, E, F;	IMPORTIN ALPHA; CHAIN: A;	KARYOPHERIN BETA2; CHAIN: B; RAN; CHAIN: C;	IMPORTIN BELA SUBUNIT; CHAIN: A; IMPORTIN ALPHA-2 SUBUNIT; CHAIN: B;	SERINE/THREONINE PROTEIN PHOSPHATASE 5; CHAIN: NULL;
SEQ FOLD score					125.55		
PMR		0.92	0.15	0.66		0.05	0.00
Verify score		0.02	-0.03	-0.20		-0,31	-0.59
Psi Blast		1.2e-46	5.1e-51	5.1e-53	1e-43	3.4e-37	0.0012
END AA		674	846	846	824		362
START AA		09	484		3	111	259
CHAIN		4	Y	V	Ø	Y	
808 B		1b3u	1004	lial	1qbk	19gr	1a17
SEQ ID NO:		273	273	273	273	273	274

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PDB annotation	STRUCTURE	LIPID BINDING PROTEIN APO-E3; LIPID TRANSPORT, LIPID TRANSPORT,	HEPARIN-BINDING,	PLASMA 2 PROTEIN, HDL,	VLDL REMARK	COMPLEX (HSP24/HSP70)	HSP70, GRPE, MOLECULAR	CHAPERONE, NOCLEOI IDE	COILED-COIL, COMPLEX	(HSP24/HSP70)	ENDOCYTOSIS/EXOCYTOSI	COMPLEX MILTINITY	ENDOCYTORIS/HYOCYTORI	SYNAPTOTAGMIN	ASSOCIATED 35 KDA	PROTEIN, P35A, THREE	HELIX BUNDLE	ZINC-BINDING PROTEIN	ZINC-BINDING PROTEIN,	XNF7, BBOX,	DEVELOPMENT, 3 MID-	CONTRO A CTUT TO DE CATERIA	TRIPLE-HELIX COLLED	COIL, CONTRACTILE	PROTEIN	CONTRACTILE PROTEIN	TRIPLE-HELIX COLLED	COIL, CONTRACTILE	PROTEIN	Ve	LIPID TRANSPORT APO A-1; ILPOPROTEIN, LIPID
Compound		APOLIPOPROTEIN E; CHAIN: A;		· .	•	NUCLEOTIDE EXCHANGE	FACTOR GRPE, CHAIN: A,	B; MOLECULAR	CHANE DIVAIN;		SYNTAXIN BINDING	PROTEIN I; CHAIN: A;	CONTACTA 16; CHAIN: B;	B. C.				NUCLEAR FACTOR XNF7;	CHAIN: NULL;			THE MAN SEVEN DEFAT	MUSCLE ALPHA-ACTININ	2; CHAIN: A;		HUMAN SKELETAL	MUSCLE ALPHA-ACTININ	2; CHAIN: A;			APOLIPOPROTBIN A-1; CHAIN: A, B, C, D;
SEQ FOLD score									,								•									59.40	•				64.70
PMF score		0.27				0.18	•		-		00:0		110	11:0	•			0.54				120	10.0			-		٠			
Verify		-0.14				-0.54					-0.00	·	0.21	; ,				-0.40				71.0	01.0								
Psi Blast		0.00017				0.00012					2.4e-06		120.07	100717				9.6e-14				1 20 07	10-27-1			1.2e-07					2.4e-10
END AA		251			•	289					268		020	2				143	•			360	007			349					427
START		131				158					143		145	}				106				1/0	740			85					224
CHAIN		Ą				A					В		•	¢								\[\bar{\}	<			A					А
PDB CI		1bz4				1dkg					[up]		1073					lfre					nnhı			Iquu					lavi
SEQ ID NO:		275				275			_		275	•	275					275				346	C/7	,		275					276

Table

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PDB annotation	CHOLESTEROL METABOLISM, 2 ATHEROSCLEROSIS, HDL, LCAT-ACTIVATION	ENDOCYTOSIS/EXOCYTOSI S SYNAPTOTAGMIN ASSOCIATED 35 KDA PROTEIN, P35A, THREE HELIX BUNDLE	BNDOCYTOSIS/EXOCYTOSI S SYNAPTOTAGMIN ASSOCIATED 35 KDA PROTEIN, P35A, THREE HELIX BUNDLE	CONTRACTILE PROTEIN TRIPLE-HELIX COILED COIL, CONTRACTILE PROTEIN	TRANSCRIPTION REGULATION SIGMA70; RNA POLYMERASE SIGMA FACTOR, TRANSCRIPTION REGULATION	TRANSCRIPTION REGULATION SIGMA70; RNA POLYMERASE SIGMA FACTOR, TRANSCRIPTION REGULATION	GENE REGULATION POZ DOMAIN; PROTEIN- PROTEIN INTERACTION DOMAIN, TRANSCRIPTIONAL 2 REPRESSOR, ZINC-FINGER PROTEIN, X-RAY CRYSTALLOGRAPHY, 3 PROTEIN STRUCTURE, PROMYELOCYTIC
Сотроипа		SYNTAXIN-1A; CHAIN: A, B, C;	SYNTAXIN-1A; CHAIN: A, B, C;	HUMAN SKELETAL MUSCLE ALPHA-ACTININ 2; CHAIN: A;	RNA POLYMERASE PRIMARY SIGMA FACTOR; CHAIN: NULL;	RNA POLYMERASE PRIMARY SIGMA FACTOR; CHAIN: NULL;	PROMYELOCYTIC LEUKEMÍA ZINC FINGER PROTEIN PLZP; CHAÍN: A;
SEQ FOLD score				65.07	77.03		
PMF score		-0.19	0.01			90.0	1.00
Verify score		0.19	0.09			-0.16	0.56
Psi Blast	·	4.8e-08	1.2e-10	1.2e-24	4.8e-09	1.2e-06	3,46-21
A END		332	358	445	469		149
START AA		228	239	081	166	291	56
CHAIN		∢	∢	V			4
PDB U		lez3	lez3	Iquu	lsig	lsig	1buo
SEQ ID NO:		276	276	276	276	276	277

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PDB annotation	LEUKEMIA, GENE REGULATION	GENE REGULATION POZ DOMAIN; PROTEIN- PROTEIN INTERACTION	DOMAIN, TRANSCRIPTIONAL 2 REPRESSOR TING-FINGER	PROTEIN, X-RAY CRYSTALLOGRAPHY, 3 PROTEIN STRUCTURE,	PROMYELOCYTIC LEUKEMIA, GENE REGULATION		***************************************	COMPLEX (ZINC FINGER/DNA) COMPLEX	ONA-BINDING	PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX	(ZINC FINGER/DNA), ZINC	PROTEIN			(ZINC FINGER/DNA), ZINC	FINGER, DNA-BINDING	PROTEIN	COMPLEX (ZINC FINGER,F)	\$
Compound		PROMYBLOCYTIC LEUKEMIA ZINC FINGER PROTEIN PLZR; CHAIN: A;				OXIDOREDUCTASE(OXYGEN(A)) GALACTOSE OXIDASE (E.C.1.1.3.9) (PH		QGSR ZINC FINGER PEPTIDE; CHAIN: A;	DUTLEA	BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER. PEPTIDE; CHAIN: A;	DUPLEX	BINDING SITE; CHAIN: B,	C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A;	DUPLEX	OLIGONUCLEOTIDE	BINDING SITE; CHAIN: B, C;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER	
SEQ FOLD score	·																			
PMF score		1.00				0.19		0.87			0.13				0.09				0.10	
Verify		0.28				0.19		0.24			0.21				-0.40				-0.02	
. Psi Blast		2.4e-22				8.5e-13.		5.1e-22			3.4e-23				6e-29	•			1e-38	
END AA		146	•		!	611		215	•	!	243				2 4 4				215	
START AA		31				299		120			148				152				119	
CHAIN		⋖						¥			¥	•			∢				O .	
20 <u>8</u>		1buo				lgof		lalh			laIh				lalh				1mey	
SEQ ID NO:		277				277		278			278			0.00	2/8		•		278	

Table

				10.24		The state of the state of
PDB annotation	PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CR YSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, F PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX
Compound	PROTEIN; CHAIN: C, F, G,	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G,	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;
SEQ FOLD score						
PMF		0.89	1.00	1.00	1.00	1.00
Verify score		0.19	0.29	0.30	0.21	0.37
Psi Blast		6.8e-41	1e-44	3.4e-46	6.8e-47	1.7e-48
END		243	271	299	327	355
START AA		147	190	218	246	274
CHAIN ID		၁	ပ	ບ	၁	υ
PDB ID		lmey	1mey	Ітеу	lmey	Imey
SEQ ID NO:		278	278	278	278	278

Table 5

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PDB annotation	(ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER,	PROTEIN-DNA	INTERACTION, PROTEIN DESIGN 2 CRYSTAI	STRICTURE COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC	FINGER/DNA) ZINC FINGER,	PROTEIN-DINA	INTERACTION, PROTEIN	STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC	FINGER/DNA) ZINC FINGER,	FROIENT-DIAM	INIEKACIJON, PROIEIN	DESIGN, 2 CKISIAL	(ZINC FINGER/DNA)	COMPLEX (ZINC	FINGER/DNA) ZINC FINGER,	PROTEIN-DNA	INTERACTION, PROTEIN	DESIGN, 2 CRYSTAL	STRUCTURE, COMPLEX	CONTRACTOR OF THE CONTRACTOR O	HINGER ONA ZINC HINGER F	PROTEIN-DNA	INTERACTION, PROTEIN	DESIGN, 2 CRYSTAL	STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC	PROTEIN-DNA
Compound	•	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;				DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER	FROIEIN; CHAIN: C, F, G;				DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER	racitan; Carani: C, F, G,				DNA; CHAIN: A, B, D, B;	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;				DNA. CHAIN. A B D B.	CONSENSITS ZINC HINGER	PROTEIN: CHAIN: C. F. G.					DNA; CHAIN: A, B, D, E;	PROTEIN; CHAIN: C, F, G;
SEQ FOLD score												•				•																	
PMF score		1.00					1.00	- *	,		•		1.00						1.00		•				8	3.5						1.00	,
Verify		0.48					0.55				•		0.31						0.32		-				0.50	7C'O						0.38	
Psi Blast		6.8e-49			-		1.7e-49	•					3.46-50						8.5e-51						1 42 50	1.46-30						1.7e-50	
END		383		_			411						439						467						307	<u> </u>						523	
START		302					330						358						386						717	‡ †						442	
CHAIN		ပ					ပ						ن ن						· U						c	 ر			,			၁	
PDB TD		1mey					lmey						1mey						1mey						1	r med						1mey	
SEQ ID		278		,			278						278						278						270	0/7						278	

Table 5

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PDB annotation	INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER,	PROTEIN-DNA INTERACTION, PROTEIN DESIGN 2 CB VST A1	STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER,	PROTEIN-DNA INTERACTION, PROTEIN	DESIGN, 2 CRYSTAL	STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC	PROTEIN-DNA	INTERACTION, PROTEIN	DESIGN, 2 CRYSTAL	SIRUCIURE, COMPLEA (ZINC FINGER/DNA)	COMPLEX (ZINC	PROTEIN-DNA	INTERACTION, PROTEIN	DESIGN, 2 CRYSTAL	STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER.	PROTEIN-DNA	INTERACTION, PROTEIN	STRUCTURE, COMPLEX	(ZUNC FUNGER/DINA)
Сотроипа		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;		•	DNA; CHAIN: A, B, D, E;	PROTEIN; CHAIN: C, F, G;			-	DNA; CHAIN: A, B, D, E;	PROTEIN; CHAIN; C, F, G;				DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G,			
SEQ FOLD score		100.48																					
PMF score					1.00				1.00					1.00					0.09			•	
Verify score					0.43				0.61					0.48					0.11				
Psi Blast		1.46-50	, -		3.4e-50				8.5e-50					1.7e-35					5.1e-37				
END A		524			551				579	,				585					172				
START		442			470				498					526					97				
CHAIN		ບ			ပ		•		ပ					ບ					ပ				
PDB ID		lmey	•	-	lmey				lmey					Imey	_				lmey		_		
SEQ ID NO:		278			278				278					278					278				

Table :

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PDB annotation	COMPLEX (ZINC FINGER/DNA) ZINC FINGER,	PROTEIN-DNA	DESIGN, 2 CRYSTAL	STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA) TFIIIA;	SS GENE; NMR, TFIIIA,	PROTEIN, DNA,	TRANSCRIPTION FACTOR,	SS KNA 2 GENE, DIVA BINDING PROTEIN, ZINC	FINGER, COMPLEX 3	(TRANSCRIPTION	REGULATION/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA) TFIIIA;	SS GENE; NMK, IFILIA,	PROTEIN, DNA,	TRANSCRIPTION FACTOR,	SS KINA 2 GEINE, DINA	BINDING PROTEIN, ZINC	FINGER, COMPLEA 3	(TRANSCRIPTION PEGIT ATTONIONA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA)	COMPLEX (TRANSCRIPTION)	REGULATION/DNA), RNA	POLYMERASE III, 2	TRANSCRIPTION	INITIATION, ZINC FINGER	PROTEIN	COMPLEX (TRANSCRIPTION)	COMPLEX (TRANSCRIPTION	REGULATION/DNA), RNA
Compound	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;			TRANSCRIPTION FACTOR	IIIA; CHAIN: A; 5S RNA	GENE; CHAIN: E, F;	:						TRANSCRIPTION FACTOR	IIIA; CHAIN: A; 5S RNA	GENE; CHAIN: E, F;						-	THIIA: CHAIN: A. D. 5S	RIBOSOMAL RNA GENE;	CHAIN: B, C, E, F;				1.		TEIIA; CHAIN: A, D; 5S RIROSOMAI RNA GRNE	CHAIN: B, C, E, F;	
SEQ FOLD score																													·			•	
PMF score	0.07				0.00									-0.02									190	5							1.00		
Verify	-0.05				-0.05									0.08									005	3	,		•		,		0.10		
Psi Blast	1.2e-07				8.5e-16								•	3.4e-15	-							•	126-32	3					· ,		1.7e-36		
END AA	144				211) 								243									308	}					•.		364		
START AA	119				120			,						148	-								148	2							219		
CHAIN	_ප				A									Ą									V	:							∢		
FDB EDB	lmcy				163						-			1463	٠.								1.5%	2							1tf6		
SEQ ID NO:	278				278						_ _			278		•							278	2		-					27.8		

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PDB annotation	POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA)	REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA)	REGULATION/DNA), RNA POLYMERASE III, 2	IKANSCKIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2	TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION	REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INTIATION, ZINC FINGER	PROTEIN COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION
Compound		TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, B, B:	(1 to 1) (2 to 1)	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, B, P:			TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE;	CHAIN: B, C, B, F;		TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;		TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE;	CHAIN: B, C, E, F;
SEQ FOLD score				104.66	·			•					
PMF score		1.00			: .		1.00			1.00	·	1.00	·
Verify score	-	-0.03				•	0.35			0.18		0.22	
Psi Blast	·	5.1e-37		1.7e-37			1.7e-37			3.4e-37	•	1.4e-35	
END		420		554			539			561		581	
START		275		386			387			415	•	443	
CHAIN		A		V			A			A		¥	
PDB U		1tf6		100	-		1466			1116		1tf6	
SEQ ID		278		278			278			278		278	

Table 5

				Harry Inches within the Land States of States	the state of twee to the state of
PDB annotation	REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INTITATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING- YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA- PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION, REGULATION/DNA) YING-YANG 1; TRANSCRIPTION FINITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-FINGER PROCEIN, DNA-FINGER PROCEIN, DNA-FINGER PROCEIN, DNA-FINGER PROCEIN, DNA-FINGER PROCEIN, DNA-FINGER PROCEIN PROCEIN RECOGNITION, 3
Compound		YY1; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YY1; CHAÏN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAÎN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR BLEMENT DNA; CHAIN: A, B;	YY1; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;
SEQ FOLD score					
PMF score		0.23	0.93	0.92	1.00
Verify	• 1	-0.01	-0.03	-0.27	0.10
Psi Blast		3.4e-25	3.4e-28	8.5e-32	6.8e-34
AA END		215	271	299	383
START AA		105	155	198	282
CHAIN		ပ	O	ပ	υ
PDB UD		lubd	1ubd	1ubd	lubd
SEQ ID NO:		278	278	278	278

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PDB annotation	COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-	YANG 1; TRANSCRIPTION	ELEMENT, YY1, ZINC 2	FINGER PROTEIN, DNA-	PROTEIN RECOGNITION, 3	COMPLEX (TRANSCRIPTION REGULATIONDNA)	COMPLEX (TRANSCRIPTION	YANG 1: TRANSCRIPTION	INITIATION, INITIATOR	ELEMENT, YY1, ZINC2	FINGER PROTEIN, DNA-	COMPLEX (TRANSCRIPTION	REGULATION/DNA)	COMPLEX (TRANSCRIPTION	YANG 1: TRANSCRIPTION	INITIATION, INITIATOR	ELEMENT, YY1, ZINC 2	FINGER PROTEIN, DNA-	PROTEIN RECOGNITION, 3	REGULATION/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA) YING-	YANG I; TRANSCRIPTION	INITIATION, INITIATOR	FINGER PROTFIN DNA-	PROTEIN RECOGNITION. 3	COMPLEX (TRANSCRIPTION	REGULATION/DNA)	COURT THE COURT
Compound		YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS P5	INITIATOR ELEMENT DNA: CHAIN: A R:					YY1; CHAIN: C; ADENO-	ASSOCIATED VIKUS PS INITIATOR RIEMENT	DNA; CHAIN: A, B;	•				YY1; CHAIN: C; ADENO-	ASSOCIATIED VIRUS PS INITIATOR ELEMENT	DNA; CHAIN: A, B;			٠		YY1; CHAIN: C; ADENO-	ASSOCIATED VIRUS P5	INITIATION FLEMENT	DNA; CHAIN: A, B;				ZINC FINGER PROTEIN	THE STREET STREET
SEQ FOLD score															83.90	- · · · · · ·									 -					,
PMF		1.00						1.00	•													1.00							0.03	
Verify score		0.30						0.32					,									0.14	-						-0.42	
Psi Blast		1.5e-34	<u>.</u>					le-34	•						3.4e-35							3.4e-35							3.6e-18	
END		411						467							524							523							245	
START AA	-	310						366				:			416							422							129	1
CHAIN		၁						ာ							— ပ	-						၁							A	
PDB		lubd			,			lubd							lubd							1ubd					-		2eli	
SEQ ID NO:		278	,					278		,					278							278							278	1

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PDB annotation	PROTEIN/DNA) FIVE- FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA-	BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE.	FINGER GLI, GLI, ZINC	FINGER, COMPLEX (DNA-	COAPI EV ONA BRIDING	PROTEINDNA) FIVE-	FINGER GLI; GLI, ZINC	FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING	PROTEIN/DNA) FIVE-	FINGER GLI; GLI, ZINC	FINGER, COMPLEX (DNA-	BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING	FROIDER/DIVA) FIVE-	FINGER COMPLEX ONA-	BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING	PROTEIN/DNA) FIVE-	FINGER GLI; GLI, ZINC	FINGER, COMPLEX (DNA-	BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING	PROTEIN/DNA) FIVE	FINGER GLI; GLI, ZINC	FINGER, COMPLEX (DNA-	BINDING PROTBIN/DNA)	COMPLEX (DNA-BINDING	PROTEIN/DNA) FIVE-	FINGER, COMPLEX (DNA-	BINDING PROTEIN/DNA)
Compound	GLII; CHAIN: A; DNA; CHAIN: C, D;		ZINC FINGER PROTEIN GLII: CHAIN: A: DNA:	CHAIN: C, D;		THIS BRIGGED DE CAPERI	GLII; CHAIN: A; DNA;	CHAIN: C, D;		ZINC FINGER PROTEIN	GLII; CHAIN: A; DNA;	CHAIN: C, D;			ZINC FINGER PROTEIN	CHAIN: OD.			ZINC FINGER PROTEIN	GLII; CHAIN: A; DNA;	CHAIN: C, D;			ZINC FINGER PROTEIN	GLII; CHAIN: A; DNA;	CHAIN: C, D;			ZINC FINGER PROTEIN	GLII; CHAIN: A; DNA;	(1 t)	
SEQ FOLD score			-												94.24								•									
PMF score			0.72			000	66.0			0.99		•							1.00				·	1.00			٠		9.			
Verify score			0.23		,	070) + 			0.07		-							0.23					0.16					0.14	,		
Psi Blast			2.40-52			3.40-37	3.40-34			6e-68					89-99				1.7e-34					3.4e-34					3.6e-71			
END			329			306	350			413					413				410					494					553			
START AA			144			8	2			246					274				282					366					386			
CHAIN			∢			4	¢			Α.					¥				Ą					∢					V			
PDB ID			2gli			2oli	11.00 m			2gli					2gli				2gli					2gli					2gli			
SEQ ID NO:	·		278			278	9 .			278					278				278	-		•		278					278			

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PDB annotation	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE- FINGER GLI, GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	ione (Grave (Grave
Compound	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	DNA-BINDING PROTEIN HUMAN ENHANCER- BINDING PROTEIN MBP-1
SEQ FOLD score					86.47		
PMF	1.00	-0.20	-0.20	1.00	·	1.00	0.36
Verify	0.29	0.15	0.05	0.21		0.39	-0.10
Psi Blast	2.4e-73	5.1e-11	1.5e-15	1.5e-23	1.4e-22	1.46-22	8.4e-24
END	581	207	239	397	538	542	484
START	442	137	161	299	457	484	431
CHAIN	¥	. V	V	V	4	V	
PDB ID	2gli	la1h	lalh	lalh		lalh	1bbo
SEQ ID NO:	278	279	279	279	279	279	279

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PDB annotation		COMPLEX (ZINC FINGER/DNA) ZINC FINGER.	PROTEIN-DNA	INTERACTION, PROTEIN DESIGN, 2 CRYSTAL	STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC	PROTEIN-DNA	INTERACTION, PROTEIN	CENIGN, 2 CRYSTAL CTOTTON OF THE	(ZINC FINGER/DNA)	COMPLEX (ZINC	FINGER/DNA) ZINC FINGER, PROTEIN-DNA	INTERACTION, PROTEIN	DESIGN, 2 CRYSTAL	STRUCTURE, COMPLEX	COMPLEX (ZINC	FINGER/DNA) ZINC FINGER,	PROTEIN-DNA	INTERACTION, PROTEIN	DESIGN, 2 CRYSTAL	STRUCTURE, COMPLEX	COMPLEX (ZINC	FINGER/DNA) ZINC FINGER,	PROTEIN-DIVA	DESIGN, 2 CRYSTAL	STRUCTURE, COMPLEX	1
Compound	MUTANT WITH CYS 11 1BBO 3 REPLACED BY ABU (C11ABU) (NMR, 60 STRUCTURES) 1BBO 4	DNA; CHAIN: A, B, D, B; CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, P, G;			DNA; CHAIN: A, B, D, E;	PROTEIN; CHAIN: C, F, G;			-	DNA; CHAIN: A, B, D, E;	PROTEIN CHAIN: C. P. G.				DNA: CHAIN: A. B. D. B.	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;				DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;			
SEQ FOLD score						·				·														•			
PMR		-0.20				0.95					1.00					1.00						1.00					
Verify score		10.0				0.14					0.48					0.57			•			0.79			•		
Psi Blast		1.7e-30				1.5e-42					3.4e-47					le-49						3.4e-48			-		
END AA		239			:	26£					425					453						480				· .	
START AA		160				298					345					372	· ·					400					
CHAIN		ပ				၁					၁				•	U						C					
PDB ID		lmey				1mey					lmey					1mey	•				-	lmey					1
SEQ ID NO:		279				279					279					279						279					

Table 5

						rest . Firstly, self arrest, except sounds, record
PDB annotation	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTHR ACTION PROTHIN	DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX	COMPLEX (TRANSCRIPTION REGULATION/DNA) TFILIA; 5S GENE; NMR, TFILIA, PROTEIN, DNA, TRANSCRIPTION FACTOR, 5S RNA 2 GENE, DNA BINDING PROTEIN, ZINC FINGER, COMPLEX 3 (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) THIIA; 5S GENE; NMR, THIIA, PROTEIN, DNA, TRIIA, TRANSCRIPTION FACTOR, 5S RNA 2 GENE, DNA BINDING PROTEIN, ZINC FINGER, COMPLEX 3
Compound	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	TRANSCRIPTION FACTOR IIIA; CHAIN: A; 5S RNA GENE; CHAIN: B, F;	TRANSCRIPTION FACTOR IIIA; CHAIN: A; 5S RNA GENE; CHAIN: E, F;
SEQ FOLD score			107.34			
PMF score	1.00			1.00	-0.20	-0.20
Verify score	0.51			0.61	0.22	0.03
Psi Blast	1.7e-48		1.7e-48	6.8e-36	1.76-08	5.1e-11
AA AA	536		537	542	200	238
START AA	456	٠	456	483	146	161
CHAIN	ن ن		U	O	∢	∢
80B E10	1mey		Imey	1mey	<u> </u>	11t3
SEQ ID NO:	279		279	279	972	279

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РDВ annotation	(TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) TFIIIA; 5S GENE; NMR, TFIIIA,	PROTEIN, DNA,	TRANSCRIPTION FACTOR,	BINDING PROTEIN, ZINC	FINGER, COMPLEX 3	(TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA)	DECIMELED (IRANSCRIPTION PRA	POI YMERASE III 2	TRANSCRIPTION	INITIATION, ZINC FINGER	PROTEIN	COMPLEX (TRANSCRIPTION	COMPLEX (TRANSCRIPTION	REGULATION/DNA), RNA	POLYMERASE III, 2	TRANSCRIPTION	INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION	REGULATION/DNA)	COMPLEX (TRANSCRIPTION	KEGULATION/DNA), KNA	POLYMERASE III, 2	INITIATION, ZINC FINGER	PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING- F	YANG 1; TRANSCRIPTION
Compound		TRANSCRIPTION FACTOR IIIA; CHAIN: A; 5S RNA GENB; CHAIN: B, F;						TFIIIA; CHAIN: A, D; 5S	RIBOSOMAL RNA GENE;	CRAIN: B, C, E, F;					TFIIIA; CHAIN: A, D; 5S	CHAIN: B. C. R. P.			•	-	TFIIIA: CHAIN: A. D: 5S	RIBOSOMAL RNA GENE;	CHAIN: B, C, E, F;		•	•		YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS P5	INITIATOR ELEMENT
SEQ FOLD score	•														115.39														
PMF score		0.03					·	1.00													1.00							1.00	
Verify score		0.05	•					0.24													0.28							0.22	
Psi Blast		1.1e-24		•			•	6.8e-35	,						8.5e-37						8.5e-37							3.4e-29	
END		456						496							537						538							425	
START		346						346			•				372						401	•						328	
CHAIN	÷	Ą						A							- ✓						V					,		၁	
PDB ID		1473						1tf6							1tf6						1166							1ubd	
SEQ ID NO:		279						279							279						279							279	

			P**	- lear roller in 1	- the same transfer
PDB annotation	INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA- PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX: (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INTITATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOMITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATIONDNA) YING-YANG 1; TRANSCRIPTION INITIATION INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING- YANG 1; TRANSCRIPTION NITIATOR INTIATION, INITIATOR ELBMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-
Compound	DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADBNO- ASSOCIATED VIRUS P5 INITATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;
SEQ FOLD score				86.73	
PMF score		1.00	1.00		00:1
Verify score		0.05	0.55		0.23
Psi Blast		1.2e-34	3.4e-34	3.4e-34	8.5e-33
END AA	·	453	508	537	536
START AA		353	408	428	436
CHAIN		U .	၁	ပ	υ
PDB ID		1ubd	1 ubd	1ubd	Jubd
SEQ ID NO:		279	279	279	279

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ation	NITTION SCRIPT A))R1, ZD	אישואים	VE-	ZINC	NONA)	NONE	ZE-	X (DNA	NONE	VE.	ZINC	X ON X	SINDIN	VE.	ZINC	X (DNA)	NONE	VE.	ZINC	X (DIV	NDIN	VE-		A (DINA)	
PDB annotation	ECOGI (TRAN ON/DIN	PTION ON PTION ON, AL	AR Y	NA) FI	I; GLI,	ROTED	(DNA-E	NA) FI	OMPLE	DNA-I	NA) FI	J; GLI,	OMPLE	DNA-I	NA) FI	I; GLI,	OMPLE ROTED	(DNA-I	NA) FI	i. GLI		DNA-F	NA) FI	i; Gi,	ROTED	-
GA.	PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATIONDNA)	TRANSCRIPTION REGULATION TRANSCRIPTION REGULATION, ADRI, ZINC	FINGER, NMR	COMPLEA (DNA-BINI PROTEIN/DNA) FIVE-	FINGER GLI; GLI, ZINC	FINGER, COMFLEA (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING	PROTEIN/DNA) FIVE- FINGER GLI: GLI. ZINC	FINGER, COMPLEX (DNA-	COMPLEX (DNA-BINDING	PROTEIN/DNA) FIVE	FINGER GLI; GLI, ZINC	FINGER, COMPLEX (DNA- RINDING PROTFIN/DNA)	COMPLEX (DNA-BINDING	PROTEIN/DNA) FIVE-	FINGER GLI; GLI, ZINC	FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING	PROTEIN/DNA) FIVE-	FINGER GLI; GLI, ZINC	FINGER, COMPLEX (DNA- PINDING PROFEIN/DNA)	COMPLEX (DNA-BINDING	PROTEIN/DNA) FIVE-	FINGER GLI; GLI, ZINC	FINGER, COMPLEA (DIVA- BINDING PROTEIN/DNA)	-
<u> </u>	S S S	E E E E	E S	38	Ē		ව්	PRO E	E S		PRC	Ē		8	PRC	Ž		ව්	PRC	Ž	Z Z		PRC		BE	1
		i	TETA.	Ä. Ä.			EIN	ζĄ;		LEIN	ΛA;			EIN	ζĄ;			NE SE	₹Ą;			TEIN	ďΑ;			
Compound		ADRI; CHAIN: NULL;	Marcaga adolica Diviz	ZINC FINGER FROTEL GLII; CHAIN: A; DNA;			ZINC FINGER PROTEIN	GLII; CHAIN: A; DNA; CHAIN: C. D:		ZINC FINGER PROTEIN	GLII; CHAIN: A; DNA;	<u></u>		ZINC FINGER PROTEIN	GLII; CHAIN: A; DNA;	<u>.</u> .		ZINC FINGER PROTEIN	GLII; CHAIN: A; DNA;	<u></u>		ZINC FINGER PROTEIN	GLII; CHAIN: A; DNA;	<u></u>		
S		I; CHA	DINICIP	CHAL	CHAIN: C, D;		FINGE	OLII; CHAIN CHAIN: C. D:	<u>.</u> :	FINGE	CHAID	CHAIN: C, D;		FINGE	CHAD	CHAIN: C, D;	•	FINGE	CHAD	CHAIN: C, D;		FINGE	CHAIN	CHAIN: C, D;		
		ADR	J. W.		CHEA		ZINC			ZINC	GLII	CHA		ZINC	GLII	CHA		ZINC	GLII	₹ <u>₩</u>		ZINC	GLII	CHA		
SEQ FOLD score														-	ı									,		
SEO							·			_				106.71							<u>.</u>	1				
PMF score		-0.18	91 0	6.19			1.00			1.00								1.00				1.00				
Verify score		0.01	900	9.0			0.28			0.04								0.57				0.53				-
Psi Blast		16-11	2 50.75	C7-2C			1.7e-30			7.2e-63				7.2e-63				6.8e-34	•			1.2e-61				-
END			+		<u>-</u>		\vdash			\dagger				T				\vdash				\dagger				_
		343	3/3	<u> </u>			452			510				510				507				537				_
START AA		271	187	70/			333	_		346				372				380				<u>\$</u>				
CHAIN			A	₹			A			A				A				A				A		_		
PDB ED		2adr	十	ng ₇			2gli			2gli	,			2gli				2gli				2gli				
SEQ ID NO:		279		C .			279			279				279				279				279				

Table:

				_	4 E 9		
PDB annotation	PROTEIN/DNA) FIVE- FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE- FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	TRANSFERASE MRNA PROCESSING, TRANSCRIPTION, RNA- BINDING, 2 PHOSPHORYLATION, NUCLEAR PROTEIN, ALTERNATIVE SPLICING 3 HELICAL FURN MOTIF, NUCLEOTIDYL TRANSFERASE CATALYTIC DOMAIN		COMPLEX (ZINC FINGER/DNA) ZINC FINGER, (PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CR YSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (TRANSCRIPTION E REGULATION/DNA) COMPLEX (TRANSCRIPTION I REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER	COMPLEX (TRANSCRIPTION)
Compound	GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	POLY(A) POLYMERASE; CHAIN: A;	والمتاريخ والمتارك وا	DNA; CHAIN: A, B, D, B; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	TFIIIA; CHAIN; A, D; 5S RIBOSOMAL RNA GENE; CHAIN; B, C, E, F;	YY1; CHAIN: C; ADENO-
SEQ FOLD score					103.07	110.63	87.81
PMF score		0.92	1.00		•		
Verify score		0.46	0.78				
Psi Blast		3.4e-30	5.10-95		1.4e-50	6.8e-38	5.16-33
END AA	·	543	421		449	446	225
START AA		436	19		367	283	116
CHAIN	•	<	₹		U .	Ą	υ υ
PDB U	 	2gli	165a		Ітеу	1,116	Iubd
SEQ ID NO:		279	282		286	286	286

Table 5

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PDB annotation	REGULATION/DNA) YING- YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA- PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE- FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (ZINC FINGER,DNA) COMPLEX (ZINC FINGER,DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC, FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC (FINGER, DNA-BINDING (PROTEIN	COMPLEX (ZINC FINGER, TRINGER, DAY OF THE PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2-CRYSTAL INTERECTURE, COMPLEX IN CZINC FINGER/DNA)
Compound	ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	QGSR ZINC FINGER PEPTIDE, CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;
SEQ FOLD score		96.09	·			
PMF score			0.33	0.43	0.12	0.60
Verify score			-0.13	0.29	-0.32	0.09
Psi Blast		6.8e-35	3.4e-21	3.6e-25	1.2e-25	6.8e-30
END AA		394	233	234	597	205
START AA		255	138	146	489	116
CHAIN		V	¥	¥	4	υ
PDB ED		2gli	laih	lalh	laih	lmey
SEQ ID NO:		286	287	287	287	287

COMPLEX (ZINC FINGER/DNA) ZINC FINGER,	PROTEIN-DNA INTERACTION, PROTEIN	AL MPI.EX				~			-				1 '	ď						u					1373		ļ
X (ZIN DNA) ZI	بر خوا اس	<u> </u>	(AN)	NCF		PROTEIN TAI	MPLEX NA)		INC FING	ROTEIN	TAL	MPLEX NA)		NC FING		ROTEIN	Mer ov	NA)		INC FILING	ROTEIN	TAL	MPLEX	NA)	NC FING		ROTEIN
$\square \supset c$	IN-UNA ICTION	1,2 CRYS	INGERAD	EX (ZINC	N-DNA	CTION, J	TURE, CC	EX (ZINC	JONA) ZI N-DNA	CTION, 1	I, 2 CRYS	TURE, CC INGER/D	EX (ZINC	/DNA) ZI	N-DNA	CITON, I	יי ביסדות מואט בי	INGER/D	EX (ZINC	N-DNA) 44	CTION, I	, 2 CRYS	URE, CO	NGEKUD	ex (zinc dina) zi	N-DNA	INTERACTION, PROTEIN
FINGE	PROLE	DESIGN	(ZINC F	COMPL	PROTE	NTER	STRUC	COMPL	PROTEI	INTERA	DESIGN	STRUC	COMPL	FINGER	PROTEI	INTERA	CTOTA	(ZINC F	COMPL	PROTEI	INTERA	DESIGN	STRUC	KINCH	COMPL	PROTE	INTERA
D, E; FINGER	5 4			D, E; FINGER	3, F, G;	•		D, B;	FINGER F. G.				D, B;	TNGER	, F, G,	•			D, E;	T C II					O, E;	, F, G;	
US ZINC	CHALIN			IN: A, B, US ZINC	CHAIN: 0			IN: A, B,	CHAIN: C				IN: A, B,	JS ZINC!	CHAIN				IN: A, B, J	CHAIN O		٠	•		IN: A, B, I JS ZINCE	CHAIN: C	
DNA; CHA CONSENS	rkoralin;			DNA; CHA CONSENS	PROTEIN;			DNA; CHA	PROTEIN:	•		-	NA; CHA	CONSENSI	ROLEIN;				ONA; CHA	ROTEIN					ONA; CHA CONSENSI	ROTEIN;	
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0.89				1.00	•			1.00										-	1.00	·					.00		
0.23				0.23				0.24						٠					0.10						0.48 8		
5.16-39				1.76-44				1.7e-46					6.8e-48			•			6.8e-48		•			,	3.4e-49		
233				261				289					290											†			1
13/				180				208					208		_				236					1	505		
بر									-		-												٠.	1			
THE S			十					1mey					1mey (\dashv						十			1
				×				87											37								1
	LIMES C. 13.16-39 0.23 0.69 DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER	CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G,	1mey C 180 261 1.7e-44 0.23 1.00 DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER	130 C 130	1.00 1.70 1.70 1.70 1.00 1.70 1.00 1.70 1.00 1.70 1.00 1.70 1.00 1.70	130 C. 130 DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER	Imey C 180 261 1.7e-44 0.23 1.00 DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER	Imey C 180 261 1.7e-44 0.23 1.00 DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G; C 208 289 1.7e-46 0.24 1.00 DNA; CHAIN: A, B, D, E; C CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G; C CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G; C C 208 289 1.7e-46 0.24 1.00 DNA; CHAIN: A, B, D, E; C C C C C C C C C	Imey C 180 261 1.7e-44 0.23 1.00 DNA; CHAIN; A, B, D, E; CONSENSUS ZINC FINGER	Image C 180 261 1.7e-44 0.23 1.00 DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G; Consensus Zinc Finger C 208 289 1.7e-46 0.24 1.00 DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G; D; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G; D; D; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G; D; D; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G; D; D; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G; D; D; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G; D; D; CONSENSUS ZINC FINGER C, C, C, C, C, C, C, C, C, C, C, C, C,	Imey C 180 261 1.7e-44 0.23 1.00 DNA; CHAIN: A, B, D, E;	Imey C 130 2.51 1.76-44 0.23 1.00 DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, P, G; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, P, G; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, P, G; CONSENSUS ZINC FINGER C 208 289 1.76-46 0.24 1.00 DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, P, G; C 208 290 6.8e-48 103.66 DNA; CHAIN: A, B, D, E; C 1 1 1 1 1 1 1 1 1	Imey C 208 290 6.86-48 103.66 DINA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER	Imey C 208 220 5.1E-35 0.25	Imey C 180 261 1.7e-44 0.23 1.00 DNA; CHAIN; A, B, D, B; CONSENSUS ZINC FINGER	Imay C 180 261 1.7e-44 0.23 1.00 DNA; CHAIN; A, B, D, B; CONSENSUS ZINC FINGER	Imay C 180 261 1.76-44 0.23 1.00 DNA; CHAIN; A, B, D, B; CONSENSUS ZINC FINGER	Imagy C 137 235 5.16-35 0.25 0	Imey C 130 261 1.7e-44 0.23 1.00 DNA; CHAIN; A, B, D, B;	Imey C 180 261 1.7e-44 0.23 1.00 DNA; CHAIN; A, B, D, B; CONSENSUS ZINC FINGER	Imey C 180 261 1.76-44 0.23 1.00 DNA; CHAIN; A, B, D, E, CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	Imey C 180 261 1.76-44 0.23 1.00 DNA; CHAIN; C.HAIN; C.F. G. CONSENSUS ZINC FINGER	Imey C 180 261 1.7e-44 0.23 1.00 DNA; CHAIN; C.P. G. G. G. G. G. G. G. G. G. G. G. G. G.	Imey C 180 261 1.7e-44 0.23 1.00 DNA; CHAIN; A, B, D, E; CONSENSUS ZINC FINGER	Imey C 180 261 1.7c-44 0.23 1.00 DINA; CHAIN; C. B. D. E. CONSENSUS ZINC FINGER PROTEIN; CHAIN; C. B. G. F. C. S. C. S. C. S. C. S. C. S. C. S. C. S. C. S. C. S. C. S. C. S. C. C. S. C. C. S. C. C. S. C. C. C. S. C. C. C. C. C. C. C. C. C. C. C. C. C.

Table

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PDB annotation	DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER,	PROTEIN-DNA INTERACTION, PROTEIN	DESIGN, 2 CRYSTAL	STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC	FINGER/DNA) ZINC FINGER, PROTEIN-DNA	INTERACTION, PROTEIN	DESIGN, 2 CRYSTAL	STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC FINGER (DNA) ZINC FINGER	PROTEIN-DNA	INTERACTION, PROTEIN	DESIGN, 2 CRYSTAL	STRUCTURE, COMPLEX	(ZINC FINGER/DNA) T	COMPLEX (ZINC	FINGER/DNA) ZINC FINGER	TATE A CHICAGO A CHICAGO	DESIGN 2 COVETAT	STRUCTURE, COMPLEX (1)		COMPLEX (ZINC	FINGER/DNA) ZINC FINGER;	PROTEIN-DNA	INTERACTION, PROTEIN	CENTICHTED COMPLEY	,	COMPLEX (ZINC FIL
Compound		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER	FKOTEIN; CHAIN: C, F, G;	•		DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER PROTEIN; CHAIN; C, F, G;					DNA; CHAIN: A, B, D, E; CONSENSIIS ZINC FINGER	PROTEIN; CHAIN: C, F, G;	•				DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER PROTEIN: CHAIN: CHO.	TOTAL CENTY C. I. C. I. C.		,	•	DNA; CHAIN: A, B, D, B;	CONSENSUS ZINC FINGER	PROTBIN; CHAIN: C, F, G,				DNA; CHAIN: A, B, D, E;
SEQ FOLD score							·																							
PMF score		1.00				1.00				-	90	0.1					ļ	1.00						1.00		-				1.00
Verify score		0.11				0.25					9	71.0		-				0.03						0.25				٠.		0.24
Psi Blast		1.4e-49				8.5e-50					1 72 50	1.76-50						5.1e-51						3.4e-51						6.8e-51
END		373				401					5	674						457		•				485						513
START AA		292				320			-		240	9						376						404						432
CHAIN	·	U				ပ					c	ر						ပ		-				ပ		,				၁
PDB ED		1mey	-			lmey					13000	rine)					,	lmey						1mey						1mey
SEQ ID NO:	·	287				287					787	0					200	787						287		•				287

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PDB annotation	FINGER, DNA, ZINC FINGER, PROTEIN-BNA, INTERACTION, PROTEIN DESIGN 2 CENTRAL	STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER,	PROTEIN-DNA	DESIGN, 2 CRYSTAL	STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC	FINGER/DNA) ZINC FINGER, PROTEIN-DNA	INTERACTION, PROTEIN	DESIGN, 2 CRYSTAL	STRUCTURE, COMPLEX	COMPLEX (ZINC	FINGER/DNA) ZINC FINGER,		EIN	DESIGN, 2 CKISIAL	(ZINC FINGER/DNA)	COMPLEX (ZINC	FINGER/DNA) ZINC FINGER	INTERACTION, PROTEIN	DESIGN, 2 CRYSTAL	STRUCTURE, COMPLEX	CAINC FINGER/DINA)	ZINC FINGER		INTERACTION, PROTEIN DESIGN, 2 CRYSTAL
Compound	CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, P, G;		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;			DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;		•	٠	DNA: CHAIN: A. B. D. E.	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;	•			DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER PROTEIN: CHAIN: C. F. G.			•	Part Curtant A P. D. D.	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;	
SEQ FOLD score																			-							
PMF			66'0				0.93					1.00						1.00					300	cen		
Verify score			-0.25				0.26					0.15				•	•	0.32					73.0	900		
Psi Blast			5.1e-46				5.1e-39					6.8e-50					•	1e-49	•				£ 15. 12	3.1e-13		,
END AA			569				297					625						653					246	2		
START			460		·		516	·				544						572					0,0	210		
CHAIN			ວ				၁					ပ						ລ					,	5		
PDB			lmey		-		Imey					1mey	,					1mey						TIECY		
SEQ ID NO:			287				287					287					_	287					200	/07_		

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PDB annotation	STRUCTURE, COMPLEX (ZINC FINGER/DNA)	ZINC FINGER TRANSCRIPTION FACTOR SP1; ZINC FINGER, TRANSCRIPTION ACTIVATION, SP1	COMPLEX (TRANSCRIPTION REGULATION/DNA) TFIIIA; 5S GENE; NMR, TFIIIA, PROTEIN, DNA, TRANSCRIPTION FACTOR, 5S RNA 2 GENE, DNA BINDING PROTEIN, ZINC FINGER, COMPLEX 3	(TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASB III, 2 TRANSCRIPTION INITIATION, ZINC FINGER T	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION/COMPLEX (TRANSCRIPTION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION COMPLEX (TRANSCRIPTION/COMPLEX (T	COMPLEX (TRANSCRIPTION) REGULATION/DNA) COMPLEX (TRANSCRIPTION/ REGULATION/DNA), RNA [1] POLYMERASE III, 2 [1] TRANSCRIPTION
Compound		SP1F2; CHAIN: NULL;	TRANSCRIPTION FACTOR IIIA; CHAIN: A; SS RNA GENE; CHAIN: B, P;		TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, B, F;
SEQ FOLD score							108.67
PMF score		0.66	0.07		0.94	0.93	
Verify score	•	-0.38	-0.50		0.07	-0.10	
Psi Blast		1.7e-08	5.1e-16		Ie-30	5.1e-35	3.4e-37
END AA		212	597		298	326	382
START AA		489	489	·	138	181	208
CHAIN	·		V		∢	4	4
PDB ID		1sp2	<u> </u>		1tf6	1116	1476
SEQ ID NO:		287	287		287	287	287

Table

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PDB annotation	INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA	POLYMERASE III, 2	TRANSCRIPTION INITIATION, ZINC FINGER	PROTEIN	COMPLEX (TRANSCRIPTION	REGULATION/DNA)	REGULATION/DNA), RNA	POLYMERASE III, 2	TRANSCRIPTION	INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION	REGULATION/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA), KNA	IKANSCRIPTION	INITIATION, ZINC FINGER	COMPLEX (TRANSCRIPTION	REGULATION/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA), RNA (I)	POLYMERASE III, 2	TRANSCRIPTION	INITIATION, ZINC FINGER	PROTEIN-	COMPLEX (TRANSCRIPTION	REGULATION/DINA)	REGIT ATTON/DNA) RNA	POLYMERASE III, 2
Compound		TFIIIA; CHAIN: A, D; SS RIBOSOMAL RNA GENE;	CHAIN: B, C, E, F;				TFIIIA; CHAIN: A, D; 5S	RIBOSOMAL RNA GENE;					TRIIIA; CHAIN: A, D; 5S	RIBOSOMAL RNA GENE;	CHAIN: B, C, E, F;				THILLY CHAIN: A D. SS	RIBOSOMAL RNA GENE:	CHAIN: B. C. E. F.						TFIIIA; CHAIN: A, D; 5S	KIBOSOMAL KNA GENE; CHAIN: B C R R.		
SEQ FOLD score																							•					:		
PMF score		96.0					1.00						06.0						0.57	}				,			0.43			
Verify		-0.04					-0.10		<u>. </u>				0.03				 		-0.35	}							-0.22			
Psi Blast		3.4e-37					8.5e-37			•			3.4e-38						10.33	3							1.26-31			
END AA		382					443						522						578	2							653			
START		237					293						377						405	}			-				489			
CHAIN		Ą					Ą						4						4	•							¥			
PDB U		1tf6					1tf6						1466				 		1#6	}							1476			
SEQ ID NO:		287					287						287						787	2							287			

INGER RIPTIO YING- PTION, 3 PTION, 3 RIPTIO FILON, 3 RIPTIO YING- PTION YIN						
Q.D. PDB CHAIN (CHAIN START END DID) Psi Norte Score Score Score Score Score Score Score 1ubd C 124 233 1.2e-21 -0.42 0.30 1ubd C 140 261 1.5e-27 0.01 1.00 1ubd C 183 289 1.7e-32 0.01 1.00 1ubd C 244 345 1.7e-33 0.22 1.00	PDB annotation	TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION TO INITIATION, INITIATOR FILEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION) REGULATION/DNA) YING- [1] YANG 1; TRANSCRIPTION ', INITIATION, INITIATOR ELEMENT, YY1, ZÎNC 2 FINGER PROTEIN, DNA- PROTEIN RECOGNITION, 3 [1] COMPLEX (TRANSCRIPTION) REGULATION/DNA)
O D PDB CHAIN START END Psi Verify PMF	Compound		YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;
Q.ID PDB CHAIN START END Psi Verify 10. D AA Blast score 1ubd C 124 233 1.2e-21 -0.42 0 1ubd C 140 261 1.5e-27 0.01 1 1ubd C 183 289 1.7e-32 0.01 1 1ubd C 244 345 1.7e-33 0.22 1	SEQ FOLD score					
QID PDB CHAIN START END Psi 100: Inbd C 124 233 1.2e-21 - 1ubd C 140 261 1.5e-27 0 - - - 1ubd C 183 289 1.7e-32 0 1ubd C 244 345 1.7e-33 0	PMF score		0.30	1.00	1.00	1.00
O.D. PDB CHAIN START END O.D. D. D. AA AA 1ubd C 124 233 1ubd C 183 289 1ubd C 244 345	Verify score		-0.42	0.01	0,01	0,22
O.D. PDB CHAIN START O.D. IND C 124 1ubd C 183 1ubd C 183	Psi Blast		1.26-21	1.5e-27	1.7e-32	1.7e-33
O'D PDB CHAIN O'D D DB Iubd C Iubd C Iubd C	AA B		233	261	289	345
O D BOB Inpp Inpp Inpp Inpp Inpp Inpp Inpp Inp	START AA		124	140	183	244
A SO	CHAIN		O	U	U	U
SEQ. ID NO: 287	PDB TD		1ubd	lubd -	1ubd	Iubd
	SEQ ID NO:		287	287	287	287

Table

PDB annotation	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING- YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA- PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING- TANG 1; TRANSCRIPTION (I) INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FUGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION) REGULATION/DNA) YING- [[] YANG 1; TRANSCRIPTION []
Compound	YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT
SEQ FOLD score				92.78	
PMF	1.00	0.99	86.0		0.95
Verify score	-0.10	000	-0.23		-0.09
Psi Blast	5.1e-35	1.4e-34	8.56-35	5.1e-36	5.1e-36
A EN	373	401	457	486	485
START AA	272	300	356	376	384
CHAIN	U	υ	ບ	ပ	ပ
PDB UD	1ubd	1ubd	Iubd	lubd	lubd
SEQ ID NO:	287	287	287	287	287

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PDB annotation	INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA- PROTEIN RECOGNITION, 3	COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING- YANG 1; TRANSCRIPTION INITIATION, INITIATOR	FINGER PROTEIN, DNA- FINGER PROTEIN, DNA- PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGIT ATTONDAA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING- YANG 1; TRANSCRIPTION	ELEMENT, YY1, ZINC2 FINGHR PROTHIN DNA.	PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION	COMPLEX (TRANSCRIPTION)	YANG 1; TRANSCRIPTION		FINGER PROTEIN, DNA-	COMPLEX (TRANSCRIPTION) REGULATIONDNA)	COMPLEX (DNA-BINDING	FROTEIN/DNA) FIVE-	¥ &	o
Compound	DNA; CHAIN: A, B;		YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;		YYI; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 INITIATOR ELEMENT	DIAN, CLEMIN: A, B,		YY1; CHAIN: C; ADENO-	ASSOCIATED VIRUS PS INITIATOR ELEMENT	DNA; CHAIN: A, B;	·.	•	ZINC FINGER PROTEIN	GLII; CHAIN: A; DNA; CHAIN: C. D;	•	ZINC FINGER PROTEIN
SEQ FOLD score		-					·				.*		·	•		92.67
PMF			0.88		0.84			00.1	-				0.80			
Verify score		-	-0.33		-0.04	- -		0.02					0.23			
Psi Blast			1.76-34		1.2e-25			1.2e-33					1.7e-26			7.2e-67
END AA	:		513		569			653					260		1.	319
START AA			412		468			552	•				137			180
CHAIN			ບ		ပ			υ υ			,		Ą			A
PDB TD			1ubd		lubd			lubd					2gli			2gli
SEQ ID NO:			287		287			287					287			287

Table 5

PDB annotation	PROTEIN/DNA) FIVE- FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE- FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA-	BINDING PROTEIN/DNA) COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE- FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE- FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE- FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	0	ی و	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE. FINGER GEJ; GLJ, ZINC FU FINGER, COMPLEX (DNA-FU BINDING PROTEIN/DNA) FU
Compound	GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLI1; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;
SEQ FOLD score		·						
PMF		0.95	0.94	0.33	0.88	0.81	0.11	0.95
Verify score		0.28	0.08	-0.03	0.03	0.04	-0.22	0.18
Psi Blast		4.8e-63	7.2e-67	3.6e-66	6.8e-33	6.8e-34	3.6e-65	2.4e-66
END		319	347	459	400	484	627	654
START AA		183	208	264	272	356	404	488
CHAIN		¥	∀	∢	⋖	₹	4	∢
PDB TD		2gli	2gli	2gli	2gli	2gli	2gli	2gli
SEQ ID NO:		287	287	287	287	287	287	287

Table 5

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PDB annotation	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE- FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)		IMMUNOGLOBULIN IMMUNOGLOBULIN, KAPPA LIGHT-CHAIN DIMER	COMPLEX	(ANTIBODY/ANTIGEN) FAB-	(ANTIBODY/ANTIGEN), ANGIOGENIC FACTOR	COMPLEX (HUMANIZED	ANTIBODY/HYDROLASE)	MURAMIDASE;	HUMANIZED ANTIBODY,	ANTIBODI COMPLEA, FV,	COMPLEX (HUMANIZED	ANTIBODY/HYDROLASE)	IMMUNE SYSTEM REIV, "U STABII IZED	IMMUNOGLOBULIN	JONES	2 PROTEIN, IMMUNE SYSTEM	ANTIBODY THERAPEUTIC, U)	ANTIBODY, CD52		war in the second		AB-IBP	¥.	STRUCTURE 2.7A [1] RESOLUTION BINDING 2 [1]
Compound	ZINC FINGER PROTBIN GLII; CHAIN: A; DNA; CHAIN: C, D;		IMMUNOGLOBULIN; CHAIN: A, B;	PAB FRAGMENT: CHAIN:	L, H, J, K; VASCULAR RNDOTHEI IAI GROWTH	FACTOR; CHAIN: V, W;	HULYS11; CHAIN: A, B, D,	E; LYSOZYME; CHAIN: C,	ř.	٠.		•		IG KAPPA CHAIN V-I REGION REI: CHAIN: A R-		•		CAMPATH-1H:LIGHT	CHAIN; CHAIN: L;	CAMPATH-IH:HEAVY	CHAIN; CHAIN: H;	PEPTIDE ANTIGEN; CHAIN: P:	IGM RF 2A2; CHAIN: A, C,	E; IGM RF 2A2; CHAÎN: B,	D, F; IMMUNOGLOBULIN G BINDING PROTEIN A;
SEQ FOLD score								•																	
PMF score	0.89		0.43	0.17			0.47							0.16		_		0.29					0.27		
Verify score	0.11		-0.33	-0.05			-0.56							-0.42				-0.00					-0.12		
Psi Blast	3.4e-32		5.1e-34	3.46-35			3.4e-34							6.8e-35				1.2e-33				-	3.4e-35		
END	652		103	103	-		103							103				103					103		
START AA	496		34	34			34							34	•			34					34		
CHAIN	¥		∢	1.1			Ą							∢				L					Ą		
PDB ID	2gli		156d	1bi1			Ibvk							1bww				lœl					1dee		
SEQ ID NO:	287		288	288			288							288				288					288		

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PDB annotation	OUTSIDE THE ANTIGEN COMBINING SITE SUPERANTIGEN FAB VH3 3 SPECIFICITY	IMMUNE SYSTEM IMMUNOGLOBULIN FOLD, ANTIBODY, IGM, FV					Hali Em v° Hali	OXIDOREDUCTASE THIOREDOXIN M, THIOREDOXIN CH2, CHLOROPLASTIC
Compound	CHAIN: G, H;	IGM MEZ IMMUNOGLOBULIN; CHAIN: L; IGM MEZ IMMUNOGLOBULIN;	CHAIN: H; IMMUNOGLOBULIN FV FRAGMENT OF A HUMANIZED VERSION OF THE ANTI-CD18 1FGV 3 ANTIBODY T52' (HUH52- AA FV) 1FGV 4	IMMUNOGLOBULIN IMMUNOGLOBULIN M (IG-M) FV FRAGMENT IIGM 3	IMMUNOGLOBULIN FAB FRAGMENT OF A HUMANIZED VERSION OF THE ANTI-CD18 2FGW 3 ANTIBODY H52 (HUH52- OZ FAB) 2FGW 4	DIHYDROFOLATE REDUCTASE (B.C.1.5.1.3) COMPLEX WITH FOLATE 1DRF 3	DIHYDROFOLATE REDUCTASE (B.C.1.5.1.3) COMPLEX WITH FOLATE 1DRF 3	CHLOROPLAST THIOREDOXIN M CH2; CHAIN: A;
SEQ FOLD score	·						281.49	
PMF		0.16	0.12	0.55	0.43	1.00		0.09
Verify score		-0.37	-0.36	-0.26	-0.15	0.81		0.19
Psi Blast		6.8e-34	1.7c-36	5.1e-34	1.4e-36	1.5e-77	1.5e-77	1.5e-22
END AA		103	103	103	103	187	187	209
START AA		¥.	34	34	34	2	3	102
CHAIN		J	L	L	T			A
PDB UD		ldql	Ifgv	ligm	2fgw	ldrf	1drf	1dby
SEQ ID NO:		288	288	288	288	289	289	290

Table 5

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PDB annotation	THIOREDOXIN	OXIDOREDUCTASE DIMER, THIOREDOXIN, X-RAY	CRYSTALLOGRAPHY,	OXIDOREDUCTASE	ELECTRON TRANSPORT	ELECTRON TRANSPORT	ELECTRON TRANSPORT ELECTRON TRANSPORT	ELECTRON TRANSPORT	ELECTRON TRANSPORT	ISOMERASE ALPHA/ALPHA-	GLUCOSAMINE 2-	EPIMERASE	ELECTRON TRANSPORT	ALPHA/BETA OPEN-	TWISTED PROTEIN, THIOL-	DISULFIDE	T7 DNA POLYMERASE, DNA	NUCLEOTIDYL 2	TRANSFERASE,	SEQUENCING, T	THIOREDOXIN,	PROCESSIVITY FACTOR, 3	COMPLEX	(HYDROLASE/ELECTRON TO THE ANSPORT/INNA)	PORT	HTRX, HCH1, CH1;		ELECTRON TRANSPORT	LO'		PEPTIDE RECOGNITION [1] PEPTIDE RECOGNITION, [1]
Compound		THIOREDOXIN; CHAIN:	1		THIOREDOXIN F; CHAIN:	A, B;	THIOREDOXIN F; CHAIN: A:	THIOREDOXIN M; CHAIN:	A, B;	N-ACYL-D-	GLUCOSAMINE 2- FPIMERASE: CHAIN: A. B:		THIOREDOXIN; CHAIN: A;	-			DNA POLYMERASE;	CHAIN: B: DNA: CHAIN: P.		,					THIOREDOXIN H; CHAIN:	NULL;			ELECTRON TRANSPORT THIOREDOXIN 2TRXA 2	C WWITZ	PSD-95; CHAIN: A; CRIPT; CHAIN: B;
SEQ FOLD score														÷																	
PMF score		90.0			-0.03		0.04	-0.06		0.42			90.0				0.04			,					-0.01				-0.07		0.37
Verify		0.07			0.28		0.10	0.46		0.25			0.35				0.73								0.08		-		0.33		-0.13
Psi Blast		3.4e-22			1.5e-20		1.5e-20	3.4e-23		2.4e-06			1.2e-26				6.8e-23								3.4e-20				3.4e-23		3.6e-07
END AA		209			207		202	210		648			206	_			509					_			210				209		83
START		103			26		26	99		528			102	•			66								88		•		26		36
CHAIN					₩.		∢	A		Ą			A				ф												∀		Ą
PDB U		lerv			1f9m		Ifaa	1fb6		1fp3			1quw				167p								1tof				2trx		1be9
SEQ ID		290			290		290	290		290			290				290								290	•			290		291

able 5

PDB annotation	PROTEIN LOCALIZATION	SIGNAL TRANSDUCTION HDLG, DHR3 DOMAIN; SIGNAL TRANSDUCTION, SH3 DOMAIN, REPEAT	OXIDOREDUCTASE BETA- FINGER	MEMBRANE PROTEIN/OXIDORBDUCTAS E BETA-FINGER, HETERODIMER	PEPTIDE RECOGNITION PSD-95; PDZ DOMAIN, NEURONAL NITRIC OXIDE SYNTHASE, NMDA RECEPTOR 2 BINDING	HYDROLASE HYDROLASB, NAD BINDING PROTEIN THE	OXIDOREDUCTASE OXIDOREDUCTASE, TROPANE ALKALOID BIOSYNTHESIS, REDUCTION OF 2 TROPINONE TO TROPINE, [7] SHORT-CHAIN DEHYDROGENASE	OXIDOREDUCTASE OXIDOREDUCTASE, TROPANE ALKALOID BIOSYNTHESIS, REDUCTION OF 2 TROPINONE TO TROPINE, FU
Compound		HUMAN DISCS LARGE PROTEIN; CHAIN: NULL;	NEURONAL NITRIC OXIDE SYNTHASE (RESIDUES 1-130); CHAIN: A:	ALPHA-I SYNTROPHIN (RESIDUES 77-171); CHAIN: A; NEURONAL NITRIC OXIDE SYNTHASE (RESIDUES I-130); CHAIN: B;	POSTSYNAPTIC DENSITY PROTEIN 95; CHAIN: A;	S. ADENOSYLHOMOCYSTEI NE HYDROLASE; CHAIN: A, B;	TROPINONE REDUCTASE- I; CHAIN: A, B;	TROPINONE REDUCTASE- I; CHAIN: A, B;
SEQ FOLD score							·	84.17
PMF score		0.99	0.45	0.1	0.41	0.01	1.00	
Verify score		0.76	0.42	1.04	0.55	-0.13	0.10	
Psi Blast		2.4e-14	3.6e-17	9.6e-17	7.2e-16	0.001	8.5e-63	8.5e-63
END		06	701	83	87	62	286	289
START AA		10	13	13	10	34	32	32
CHAIN			¥	¥	∢	¥	₹	¥
PDB CI		1pdr	Igau	Iqav	1qlc	1a 7 a	lael	lael
SEQ ID NO:		291	291	291	291	294	294	294

Table 5

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PDB annotation	SHORT-CHAIN DEHYDROGENASE	OXIDOREDUCTASE OXIDOREDUCTASE, TROPANE ALKALOID BIOSYNTHESIS.	REDUCTION OF 2 TROPINONE TO TROPINE, SHORT-CHAIN DEHYDROGENASE	OXIDOREDUCTASE, OXIDOREDUCTASE,	BIOSYNTHESIS,	REDUCTION OF 2 TROPINONE TO TROPINE,	SHORT-CHAIN DEHYDROGENASE	OXIDOREDUCTASE NAD-	OXIDOREDUCTASE, SHORT-	CHAIN ALCOHOL 2	DEGRADATION	OXIDOREDUCTASE NAD-	OXIDOREDUCTASE, SHORT	CHAIN ALCOHOL 2	DEGRADATION	OXIDOREDUCTASE SHORT	OXIDOREDUCTASE	OXIDOREDUCTASE SHORT.	OXIDOREDUCTASE	OXIDOREDUCTASE INHA; THE	
Compound		TROPINONE REDUCTASE- I; CHAIN: A, B;		TROPINONE REDUCTASE. I; CHAIN: A, B;				CIS-BIPHENYL-2,3-	DEHYDROGENASE;	CHAIN: NULL;		CIS-BIPHENYI2,3-	DEHYDROGENASE;	CHAIN: NULL;		CARBONYL REDUCTASE; CHAIN: A B C D:		CARBONYL REDUCTASE; CHAIN: A. B. C. D:		ENOYL-ACYL CARRIER PROTFIN (ACP)	REDUCTASE; 1ENY 4
SEQ FOLD score				89.11				78.30										84.06		21.90	
PMF score		1.00										1.00				1.00					
Verify score		0.16		·						•		0.43	٠			0.17				·	
Psi Blast		1.7e-65		1.7e-65				1.7e-48				1.7c-48				1.7e-62		1.7e-62		6.8e-19	
END AA	i	286		289				308				289				287		287		293	
START AA		32		32		·		33				35				33		33		32	
CHAIN	·	Я		В												Ą		∢			
PDB ID		lael		lael .	_			1bdb				1bdb				lcyd		lcyd		1eny	
SEQ ID NO:		294		294				294				294				294		294		294	

Table

PDB annotation		DEHYDROGENASE DEHYDROGENASE, 17- BETA-HYDROXYSTEROID	DEHYDROGENASE DEHYDROGENASE, 17- BETA-HYDROXYSTEROID	OXIDOREDUCTASE SHORT-CHAIN DEHYDROGENASE/REDUCT ASE, BILE ACID CATABOLISM	OXIDOREDUCTASE SHORT- CHAIN DEHYDROGENASEREDUCT ASE, BILE ACID CATABOLISM		/US02/0	OXDOREDUCTASE SEPIAPTERIN REDUCTASE TETRAHYDROBIOPTERIN OXDOREDUCTASE
Compound	CHAIN: NULL; 1ENY 5	17-BETA- HYDROXYSTEROID- DEHYDROGENASE; CHAIN: NULL;	17-BETA- HYDROXYSTEROID- DEHYDROGENASE; CHAIN: NULL;	7 ALPHA- HYDROXYSTEROID DEHYDROGENASE; CHAIN: A, B;	7 ALPHA- HYDROXYSTEROID DEHYDROGENASE; CHAIN: A, B;	OXIDOREDUCTASE 3- ALPHA, 20-BETA- HYDROXYSTEROID DEHYDROGENASE (E.C.1.11.53) 1HDC 3 COMPLEXED WITH CARBENOXOLONE 1HDC 4	OXIDOREDUCTASE 3- ALPHA, 20-BETA- HYDROXYSTEROID DEHYDROGENASE (E.C.1.11.53) 1HDC 3 COMPLEXED WITH CARBENOXOLONE 1HDC 4	SEPIAPTERIN REDUCTASE; CHAIN: NULL;
SEQ FOLD score		112.71		84.13			84.15	58.34
PMF score			1.00		1.00	0.94		
Verify score			0.31		0.28	0.31		
Psi Blast		1.7e-47	1.7e-47	le-64	1e-64	1.7e-69	1.7e-69	1.2e-32
END AA		296	269	306	264	284	236	284
START		36	36	29	33	33	33	32
CHAIN				¥	₹	V	∢	
PDB U		lfds	lfds	Ifmc	1ffic	Ihdc	Ihdc	loss
SEQ ID NO:		294	294	294	294	294	294	294

					•							
PDB annotation	OXIDOREDUCTASE SEPIAPTERIN REDUCTASE, TETRAHYDROBIOPTERIN, OXIDOREDUCTASE	OXIDOREDUCTASE NAPHTHOL REDUCTASE; OXIDOREDUCTASE	OXIDOREDUCTASE NAPHTHOL REDUCTASE; OXIDOREDUCTASE	OXIDOREDUCTASE OXIDOREDUCTASE, TROPANE ALKALOID	BIOSYNTHESIS, REDUCTION OF 2 TROPINONE TO	PSEUDOTROPINE, SHORT- CHAIN DEHYDROGENASE	OXIDOREDUCTASE, OXIDOREDUCTASE,	TROPANE ALKALOID BIOSYNTHESIS,	TROPINONE TO TO PSEUDOTROPINE, SHORT-	T >USD	.	CELL ADHESION PROTEIN ; A-DOMAIN INTEGRIN, CELE ADHESION PROTEIN, H GLYCOPROTEIN, III EXTRACELLULAR 2 MATRIX, CYTOSKELETON,
Compound	SEPIAPTERIN REDUCTASE; CHAIN: NULL;	TRIHYDROXYNAPHTHAL ENE REDUCTASE; CHAIN: A, B;	TRIHYDROXYNAPHTHAL ENE REDUCTASE; CHAIN: A. B;	TROPINONE REDUCTASE- II; CHAIN: A, B;			TROPINONE REDUCTASE- II; CHAIN: A, B;			OXIDOREDUCTASE (FLAVOENZYME) GLUTATHIONE REDUCTASE (B.C.1.6.4.2), OXIDIZED FORM (E) 3GRS 4		INTEGRIN; CHAIN: NULL;
SEQ FOLD score		78.32		95.13		·			• .			
PMF	0.94		1.00				00'1			0.07		0.95
Verify score	0.27		0.58				0.25			-0.75		0.16
Psi Blast	1.2e-32	1.7e-60	1.7e-60	3.4e-61			3.4e-61			0.0085		1.2e-09
END	225	284	797	301			286			29		136
START AA	36	25	34	31			32			14		7
CHAIN ID		Ą	Ą	Ą			¥					·
PDB CI	loaa	1ybv	1ybv	2ae2			2ac2			3grs		Lido
SEQ ID NO:	294	294	294	294			294			294		295

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PDB annotation	CELL ADHESION LFA-1, ALPHA-L\BETA-2 INTEGRIN, A-DOMAIN; ILFA 8	COMPLEX (TRANSCRIPTION FACTOR/DNA)	TRANSCRIPTION FACTOR, PROTEIN-DNA COMPLEX, CYTOKINE 2 ACTIVATION, COMPLEX (TRANSCRIPTION, HACTOR (DNA)	DNA-BINDING HMGA DNA- BINDING HMG-BOX DOMAIN A OF RAT HMGI;	DNA-BINDING HMGA DNA- BINDING HMG-BOX DOMAIN A OFRAT HMGI:	1AAB 8 HMG-BOX 1AAB 20	DNA BINDING PROTEIN HMG BOX, DNA BENDING, DNA RECOGNITION, CHROMATIN, NMR, DNA 2	TEIN NDING, N, DNA 2	GENE REGULATIONDNA MI HMG-1, AMPHOTERIN, MEPARIN-BINDING PROTEIN P30; HIGH-MOBILITY GROUP DOMAIN-BENT DNA, PROTEIN-DRUGHDNA 2 COMPLEX, GENE MEGULATIONDNA
Compound	CDIIA; ILFA 5 CHAIN: A, B; ILFA 6	STAT3B; CHAIN: A; 18- MER	DESOXYOLIGONUCLEOTI DE; CHAÎN: B;	HIGH MOBILITY GROUP PROTEIN; 1AAB 5 CHAIN: NULL; 1AAB 6	HIGH MOBILITY GROUP PROTEIN; 1AAB 5 CHAIN:		NON HISTONE PROTEIN 6 A; CHAIN: A;	NON HISTONE PROTEIN 6 A; CHAIN: A;	HIGH MOBILITY GROUP 1 PROTEIN; CHAIN: A; DNA (5-D(*CP*CP*(IDO) CHAIN: B; DNA (5- CHAIN: C;
SEQ FOLD score				·			54.43		
PMF	0.07	10:0		1.00	0.99			1.00	66:0
Verify score	0.10	0.02		0.32	0.67	***	·	0.52	0.63
Psi Blast	1.2e-06	3.4e-05		4.8e-14	8.5e-05		6.8e-15	6.8e-15	0.00014
END	142	339		294	298		317	299	. 298
START AA	2	173		239	249		224	249	249
CHAIN	Ą	A					¥	, V	∢
PDB	llfa	lbgl	·	laab	1aab		1cg7	lcg7	lckt
SEQ ID NO:	295	298		300	300		300	300	300

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PDB annotation			COMPLEX (DNA-BINDING PROTEIN/DNA)		GENE REGULATIONDNA 'A HMG-D; PROTEIN-DNA COMPLEX, HMG DOMAIN, (f) NON-SEQUENCE SPECIFIC CHROMOSOMAL PROTEINED HMG-D	GROWTH FACTOR/GROWTH FACTOR RECEPTOR FGP2; LA FGPR2; IMMUNOGLOBULLY (G)LIKE DOMAINS BELONGING TO THE L-SET
Compound	DNA-BINDING HIGH MOBILITY GROUP PROTEIN FRAGMENT-B (HMGB) (DNA-BINDING 1HME 3 HMG-BOX DOMAIN B OF RAT HMG1) (NMR, 1 STRUCTURE)	DNA-BINDING HIGH MOBILITY GROUP PROTEIN FRAGMENT-B (HMGB) (DNA-BINDING 1HME 3 HMG-BOX DOMAIN B OF RAT HMG1) (NMR, 1 STRUCTURE) 1HME 4	HUMAN SRY; 1HRY 6 CHAIN: A; 1HRY 7 DNA; 1HRY 9 CHAIN: B; 1HRY 10	DNA-BINDING HIGH MOBILITY GROUP PROTEIN 1 (HMG1) BOX 2, COMPLEXED WITH 1HSM 3 MERCAPTOETHANOL (NMR, MINIMIZED AVERAGE STRUCTURE) 1HSM 4	DNA (5'- D(*GP*CP*GP*AP*TP*AP* TP*CP*GP*C)-3'); CHAIN: C, D; HIGH MOBILITY GROUP PROTEIN D; CHAIN: A, B;	FIBROBLAST GROWTH FACTOR 2; CHAIN: A, B, C, D; FIBROBLAST GROWTH FACTOR RECEPTOR 2; CHAIN: B, F, G, H;
SEQ FOLD score						
PMF score	0.88	06:0	0.16	0.41	0.87	0.11
Verify score	0.34	0.36	-0.08	0.10	0.51	-0.07
Psi Blast	2.4e-15	1e-09	1.7e-05	5.1e-10	1.7e-05	7.2e-07
END AA	292	298	299	319	298	669
START AA	240	250	249	250	249	613
CHAIN			A		¥	Ð
PDB UI	lhme	Ihme	lhry	Ihsm	lqrv	lev2
SEQ ID NO:	300	300	300	300	300	301

		_								·
PDB annotation	SUBGROUP WITHIN IG-LIKE DOMAINS, B-TREFOIL FOLD			LIGASE CBL, UBCH7, ZAP- 70, E2, UBIQUITIN, E3, PHOSPHORYLATION, 2 TYROSINE KINASE, UBIQUITINATION, PROTEIN	DEGRADATION,	METAL BINDING PROTEIN RING FINGER PROTEIN MAT1; RING FINGER (C3HC4)	DNA-BINDING PROTEIN V(D)J RECOMBINATION ACTIVATING PROTEIN 1;	RAG1, V(D)I RECOMBINATION, ANTIBODY, MAD, RING FINGER, 2 ZINC BINUCLEAR CLUSTER, ZINC FINGER, DNA-BINDING PROTEIN	DNA INTEGRATION DNA CONTEGRATION, AIDS, AID FOLYPROTEIN, AIDS HYDROLASE, 2 ENDONUCLEASE, POLYNUCLEOTIDYL TRANSFERASE, DNA CONTEGRASE, CONTEGRA	DNA INTEGRATION DNA III INTEGRATION, AIDS, III POLYPROTEIN, III
Compound			VIRUS EQUINE HERPES VIRUS-1 (C3HC4, OR RING DOMAIN) 1CHC 3 (NMR, 1 STRUCTURE) 1CHC 4	SIGNAL TRANSDUCTION PROTEIN CBL; CHAIN: A; ZAP-70 PEPTIDE; CHAIN: B; UBIQUITIN- CONTUGATING ENZYME	B12-18 KDA UBCH7; CHAIN: C;	CDK-ACTIVATING KINASE ASSEMBLY FACTOR MAT1; CHAIN: A;	RAGI; CHAIN: NULL;		HIV-1 INTEGRASE; CHAIN: NULL;	INTBGRASE; CHAIN: A, B, C;
SEQ FOLD score										
PMF score			0.83	0.52		0.27	0.65		0.22	0.06
Verify			0.02	-0.34		0.30	0.03		06'0-	-0.63
Psi Blast			1.4e-12	1.40-12		1.5e-05	3.4e-16		6.8e-13	le-13
END			83	26		68	117		49	52
START AA			37	39		35	39		7	2
CHAIN				4		¥				ာ ပ
PDB UD			Ichc	1fbv		1825	lrmd		15hl	1613
SEQ ID			302	302		302	302		304	304

Table

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PDB annotation	HYDROLASE, 2 ENDONUCLEASE,	TRANSFERASE, DNA	BINDING 3 (VIRAL)	TRANSFERASE INTEGRASE,	ROUS SARCOMA VIRUS,	HIV, X-RAY	CKISI ALLOGKAPHI, 2	TRANSFERASE	VIRUS/VIRAL PROTEIN	INTEGRASE, ROUS	SARCOMA VIRUS, HIV, X-	RAY CRYSTALLOGRAPHY,	2 VIRUS/VIRAL PROTEIN	VIRUS/VIRAL PROTEIN SH3-	LIKE DOMAIN,	NONSPECIFIC DNA	BINDING BETA SHEET, CIS-	2 PROLINE	DNA-BINDING PROTEIN	DNA-BINDING PROTEIN,	AIDS, POLYPROTEIN	HYDROLASE DNA	INTEGRATION, INTEGRASE,	HIV, HYDROLASE,	ASPARTYL 2 PROTEASE,	ENDONUCLEASE	W.	MYOSIN MYOSIN MOTOR		k rove			MYOSIN MYOSIN MOTOR	2	
Compound				INTEGRASE; CHAIN: A, B,	C, D;				RSV INTEGRASE; CHAIN:	A, B;				INTEGRASE; CHAIN: A, B;	. •				HIV-1 INTEGRASE; CHAIN:	NULL	The second secon	HIV-1 INTEGRASE; CHAIN:	A, B, C;					MYOSIN HEAVY CHAIN;	REGULATORY LIGHT	CHAIN; CHAIN: Y;	MYOSIN ESSENTIAL	LIGHT CHAIN; CHAIN: Z;	MYOSIN HEAVY CHAIN;	CHAIN: A; M TOSIN REGULATORY LIGHT	
SEQ FOLD score																												413.91							
PMF				0.81					0.70					0.62					0.48			0.15											1.00		
Verify score				0.02			1.040	····	-0.28					-0.18					0,11			-0.88											0.50		
Psi Blast				8.5e-19					6.8e-16					5.1e-33					1.26-18			8.5e-13						0		•	•		0		·
AA A				22					16					96					96			25						803					809		
START AA				28					18					2					51			7											-		
CHAIN	·			٧					В					٧					Α			¥						∢					4		
PDB UI				1c0m					lcla					lex4					lihv			Iqs4						167 1	_				167t		-
SEQ ID NO:				304					304					304					304			304						305					305		

Table 5

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PDB annotation		MUSCLE PROTEIN MDE; MUSCLE PROTEIN	MUSCLE PROTEIN MDE; MUSCLE PROTEIN	MUSCLE PROTEIN MUSCLE PROTEIN	MUSCLE PROTEIN MUSCLE PROTEIN	CONTRACTILE PROTEIN MYOSIN MOTOR, CONFORMATIONAL CHANGES	CONTRACTILE PROTEIN MYOSIN, DICTYOSTELIUM, MOTOR, MANT, ATPASE, ACTIN-BINDING, 2 COLLED COIL	CONTRACTILE PROTEIN MYOSIN, DICTYOSTELIUM, MOTOR, MANT, ATPASE, ACTIN-BINDING, 2 COILED TO	CONTRACTILE PROTEIN ATPASE, MYOSIN, COILED COIL, ACTIN-BINDING, ATP BINDING, 2 HEPTAD REPEAT PATTERN, METHYLATION, ALKYLATION, PHOSPHORYLATION, CONTRACTILE PROTEIN	CONTRACTILE PROTEIN ATPASE, MYOSIN, COILED 中COIL, ACTIN-BINDING, ATFU BINDING, 2 HBPTAD 開 REPEAT PATTERN, 图
Compound	CHAIN; CHAIN: Y; MYOSIN ESSENTIAL LIGHT CHAIN; CHAIN: Z;	MYOSIN; CHAIN: A, B, C, D, E, F, G, H;	MYOSIN; CHAIN: A, B, C, D, E, F, G, H;	MYOSIN; CHAIN: A, B, C, D, E, F;	MYOSIN; CHAIN: A, B, C, D, E, F;	MYOSIN HEAD; CHAIN: A; MYOSIN HEAD; CHAIN: Y; MYOSIN HEAD; CHAIN: Z;	MYOSIN; CHAIN: NULL;	MYOSIN; CHAIN: NULL;	MYOSIN; CHAIN: NULL;	MYOSIN; CHAIN: NULL;
SEQ FOLD score			451.74		417.00		420.64		370.75	
PMF score	·	1.00		1.00		1.00		1.00		1.00
Verify score		0.56	·	0.57		0.45		0.29		0.71
Psi Blast		0	0	0	0	0	0	0	0	0
A A		775	111	746	746	808	709	746	644	644
START AA		1		1	1	-	2	က	2	3
CHAIN		¥	Α	¥	Ą	· V				·
PDB U		1br1	1br1	1br2	1br2	ldfk	IIvk	llvk	1mnd	1mnd
SEQ ID NO:		305	305	305	305	305	305	305	305	305

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				П		Juli I		2/012	
PDB annotation	METHYLATION, ALKYLATION, 3 PHOSPHORYLATION, CONTRACTILE PROTEIN	MUSCLE PROTEIN MUSCLE PROTEIN, MYOSIN SUBFRAGMENT-1, MYOSIN HEAD, 2 MOTOR PROTEIN	MUSCLE PROTEIN MUSCLE PROTEIN, MYOSIN SUBFRAGMENT-1, MYOSIN HEAD, 2 MOTOR PROTEIN		COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX
Compound		MYOSIN; CHAIN: A, B, C;	MYOSIN; CHAIN: A, B, C;		QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A;
SEQ FOLD score			319.94		87.27				
PMF score		1.00			·	0.29	0.86	0.25	0.25
Verify score		60.0				-0.19	0.46	0.05	0.29
Psi Blast		0	0		5.1e-29	1.7e-20	5.1e-29	8.5e-24	3.4e-25
END	·	785	807		202	116	229	244	356
START. AA			1		120	12	148	9/1	278
CHAIN		Y	Ą		V	V	V	∢	A
PDB ID		2mys	2mys		lalh	laih	laih	lalh	1alh
SEQ ID NO:		305	305		306	306	306	306	306

Table 5

PDB annotation	(ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN		CONTRACTILE LIM DOMAIN, CRP, NMR, MUSCLE DIFFERENTIATION, CONTRACTILE	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2_CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER, FINGERDA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX CINC FINGER/DNA)	COMPLEX (ZINC FINGER FINGERDANA) ZINC FINGERDONA) FINCTEIN-DNA INTERACTION, PROTEIN HOESIGN, 2 CRYSTAL IN STRUCTURE, COMPLEX IN CZINC FINGER/DNA)
Compound	DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	TRANSCRIPTION REGULATION YBAST TRANSCRIPTION FACTOR ADRI (RESIDUES 102 - 130) 1ARD 3 (AMINO TERMINAL ZINC FINGER DOMAIN) (NMR, 10 STRUCTURES) 1ARD 4 (ADRIB) 1ARD 5	CRP1; CHAIN: A;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, B; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;
SEQ FOLD score						
PMF score		0.70	0.09	1.00	1.00	0.74
Verify		80'0	-0.34	0.46	0.31	-0.34
Psi Blast		5.1e-05	9.6e-05	1e-50	3.4e-48	1.7e-36
END		203	127	200		116
START AA		176	30	119	147	17
CHAIN			4	U	ပ	U
PDB CI		lard	158t	Imey	1mey	Ітеу
SEQ ID NO:		306	306	306	306	306

		.		party plant rapps	to these species and the street	an Surrename
PDB annotation	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX	COMPLEX (ZINC FINGER, PROTEIN-DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX CZINC FINGER/DNA)	COMPLEX (ZINC FINGERDNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CR YSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX	COMPLEX (ZINC FINGER/DNA) ZINC FINGER I PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CR YSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	
Compound	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGHR PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, B; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN; A, B, D, B; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	TRANSCRIPTION REGULATION YEAST TRANSCRIPTION FACTOR ADR1 (RESIDUES 130 - 159)
SEQ FOLD score					111.82	
PMF score	0.60	0.12	1.00	0.1	·	0.60
Verify score	0.42	0.27	0.43	0.29		-0.41
Psi Blast	5.1e-41	3.4e-30	1.7e-48	le-50	1e-50	6.8e-05.
END A	356	382	4 ·	172	173	202
START AA	278	314	99		16	176
CHAIN	U	ပ	ပ	υ ,	ပ	·
PDB U	1теу	Imey	lmey	1mey	1mey	Ipaa
SEQ ID NO:	306	306	306	306	306	306

Table

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PDB annotation			ZINC FINGER TRANSCRIPTION FACTOR SP1; ZINC FINGER, TRANSCRIPTION ACTIVATION, SP1	CTOR,	(TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION/REGULATION/DNA) REGULATION/DNA) TFIIIA;	PROTEIN, DNA, TRANSCRIPTION FACTOR, SS RNA 2 GENE, DNA BINDING PROTEIN, ZINC FINGER, COMPLEX 3 (TRANSCRIPTION REGULATIONDNA)	COMPLEX (TRANSCRIPTION) REGULATION/DNA) TFILLAN SS GENE; NMR, TFILLA, FL
Compound	IPAA 3 (PAPA - CARBOXY TERMINAL ZINC FINGER DOMAIN) MUTANT WITH IPAA 4 PRO 131 REPLACED BY ALA, PRO 133 REPLACED BY ALA,	CYS 140 IPAA 5 REPLACED BY ALA (P131A,P133A,C140A) (NMR, 10 STRUCTURES) 1PAA 6	SPIF2; CHAIN: NULL;	TRANSCRIPTION FACTOR IIIA; CHAIN: A; 53 RNA GENE; CHAIN: B, F;	TRANSCRIPTION FACTOR IIIA; CHAIN: A; SS RNA	devis, cream; E, F;	TRANSCRIPTION FACTOR IIIA; CHAIN: A; 5S RNA GENE; CHAIN: E, F;
SEQ FOLD score							
PMF score			0.94	-0.09	0.25		-0.02
Verify score			0.05	0.20	-0.07		0.20
Psi Blast			1.7e-07	8.5e-16	1.5e-13		1.2e-15
END AA			203	237	116		355
START AA	•		176	176	26		278
CHAIN				∢	Ą		Ą
PDB 10			1sp2	143	Itf3		1143
SEQ ID NO:			306	306	306		306

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PDB annotation	PROTEIN, DNA, TRANSCRIPTION FACTOR, 5S RNA 2 GENE, DNA BINDING PROTEIN, ZINC FINGER, COMPLEX 3	(TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) TFIIIA; SS GENE: NMR. TFIIIA.	PROTEIN, DNA, TRANSCRIPTION FACTOR,	SS RNA 2 GENE, DNA BINDING PROTEIN, ZINC	FINGER, COMPLEX 3 (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION	POLYMERASE III, 2	TRANSCRIPTION	INITIATION, ZINC FINGER TO PROTEIN	COMPLEX (TRANSCRIPTION REGIL ATION/DNA)	COMPLEX (TRANSCRIPTIO	REGULATION/DNA), KNA ≒ POLYMERASE III, 2	TRANSCRIPTION C	PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA [[]]	POLYMERASE III, 2 [I] TRANSCRIPTION [I]
Compound			TRANSCRIPTION FACTOR IIIA; CHAIN; A; 5S RNA GENE: CHAIN; E. F.				TFIIIA; CHAÎN: A, D; 5S RIBOSOMAL RNA GENE:	CHAIN: B, C, E, F;			·	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE:	CHAIN: B, C, E, F;				TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE;	CHAIN: B, C, E, F;	
SEQ FOLD score			86.09																
PMF score							-0.06				r	0.75					0.12		
Verify		· .					0.15					0.09				·	0.37		
Pst Blast			1.5e-13				1.5e-27					1.4e-30					6.8e-17		
AA END			147				348					181				•	380		
START AA			63				176					26					278		
CHAIN			A				Ą					₹					¥		
PDB ID			EJH1	_			146		•			1tf6					1tf6		
SEQ ID NO:	·		908				306					306					306		

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acton	HINGE	SCRIPT A)	SCRIF1 (A), RNA	7	FINGE	Tata	A)	SCRIPT	A), RN/	7		HINGE		SCRIPT (A)	SCRIPT	IA), RN/	. 7		, ringe	SCRIPT	A) YIN		AICK	Z Z Z	LINA-	Teller		SCRIPT	IA) YIN	KILLIO
PDB annotation	N, ZINC	(TRAN ION/DN	NOWOI	ASE III, IPTION	N, ZINC	N V GLID	NOWO	(TRAN	NOV/DI	ASE III,	PTION	N, ZINC			TRAN	ION/DI	ASE III	NOLLA	יוא, בווער	TRAN	ION/OI	KANSC	N, INII	XXI,Z	KO LELLON		NOVNOL	(TRAN	NONOL	KANSC
Q	INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATIONDNA)	REGULATION/DNA), RNA	POLYMERASE III, 2 TRANSCRIPTION	INITIATION, ZINC FINGER	PROTEIN	REGULATION/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA), RNA	POLYMERASE III, 2	TRANSCRIPTION	INITIATION, ZINC FINGER	PROTEIN	COMPLEX (TRANSCRIPTION DECITE ATTOM/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA), RNA	POLYMERASE III, 2	TRANSCRIPTION	INTITATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION	REGULATION/DNA) YING-	YANG I; TRANSCRIPTION		ELEMENT, YYI, ZINCZ	FINGER FROIEIN, DINA-	VILLE LOS	REGIL ATION/DNA)	COMPLEX (TRANSCRIPTION	REGULATIONDNA) YING-[[]	YANG I; TRANSCRIPTION FI
	ZI ZI		2 Z	<u> </u>	召	E			2	<u> </u>		3	ă.			8	- F	<u> </u>			2	7 2	<u> </u>	코 F	T F	בן כ	<u>څ</u> ک	T	2:	¥ £
pı		D; 5S A GENE				2	A GENE		•					, D; 5S						NDENO-	RUS P5	TENT	ë.					DENO	RUS P5	MEINT.
Compound		IAIN: A	֡֝֝֝֝֡֝֝֟֝֝֓֞֝֞֝֞֝֞֝֞֝֞֝֞֝֞֝֞֝֞֝֡֝֞֝֓֞֝֞֡֝֡֝֡֡֝֡֝֡֝֡֡֝֡֡֝֡֡֝֡֡֝֡֡֝֡֝֡֡֝֡֡֝֡֡֡֡֡֡			7 7 7 7	ALM: A	C.B.F						HAIN: A	C F F					S.C.	TED VI	R ELEN	AIN: A					KIN. C. 1	TED VI	RELEN
0		TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE;	CHAIN: B, C, E, F;				TFILIA; CHALIN: A, D; 38 RTBOSOMAL RNA GENE:	CHAIN: B, C, E, F;						TEIIA; CHAIN: A, D; 5S	CHAIN: B. C. E. F.					YY1; CHAIN: C; ADENO-	ASSOCIATED VIRUS P5	INITIATOR ELEMENT	DNA; CHAIN: A, B;				٠.	YY1; CHAIN: C; ADENO-	ASSOCIATED VIRUS P5	INITIATOR ELEMENT
CΩ		E 25	ブ 			+	7 2	<u></u> 5					-	E 6	2 5	·	•			<u>\\ \</u>	<u> </u>	4	<u> </u>		<u></u>			\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\	< 1	41
SEQ FOLD score		108.72						•				•					•					. •								
PMF score			-				3.							0.98						0.94								-0.14		
Verify score							0.27		:					0.03						0.36							-	0.16		
Psi Blast		8.5e-37				50	8.26-37							5.1e-36						1.5e-32								1.4e-26		
END AA		224				1	210			•				238						229								302		
START		28				ļ	8							92	•					127								155		
CHAIN		A					∢							Ą						U								U)	
EDB CD		146				,	100							1tf6						1ubd								lubd		
SEQ ID	,	10												9	_					2							-	9		
15 ~	[306					<u>8</u>							306						306								306	}	

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PDB annotation	ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA- PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRETION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING- YANG 1; TRANSCRIPTION INTIATION, INTIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA- PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING- YANG 1; TRANSCRIPTION INTIATION, INTIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA- PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING- TYANG 1; TRANSCRIPTION INTIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA- PROTEIN RECOGNITION, 3 (I) REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-CAND 1; TRANSCRIPTION HINTIATION, INITIATOR ELEMENT, YY1, ZINC 2 FUNGER PROTEIN, DNA-FUNGER PROCENITION, 3 FU
Compound		YYI; CHAIN: C; ADENO-ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YY1; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADBNO-ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;
SEQ FOLD score				103.37	
PMF		0.07	0.53		1.00
Verify		-0.22	0.07		0.32
Psi Blast		1.7e-20	8.5e-23	1.7e-34	5.1e-34
END AA		116	356	173	172
START AA		17	280	29	71
CHAIN		၁	ပ	U	υ
EDB ED		lubd	lubd	Iubd	lubd
SEQ ID NO:	·	306	306	306	

Table 5

PDB annotation	COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-	YANG 1; TRANSCRIPTION	INITIATION, INITIATOR	ELEMENT, YY1, ZINC 2	FINGER PROTEIN, DNA-	PROTEIN RECOGNITION, 3	COMPLEX (TRANSCRIPTION REGIT ATTON/DNA)	TRANSCRIPTION	REGULATION	TRANSCRIPTION	REGULATION, ADRI, ZINC	FINGER, NMK	TRANSCRIPTION	TRANSCRIPTION	REGULATION, ADRI, ZINC	FINGER, NMR	TRANSCRIPTION	REGULATION	TRANSCRIPTION	REGULATION, ADRI, ZINC	FINGER, NMR	TRANSCRIPTION	REGULATION	IKANSCKIPTION THE	, AUKI, ZINC	TE ANSCRIPTION FILE	ALICA MININA MIN	NC	OR1. ZINC	FINGER, NMR	COMPLEX (DNA-BINDING 14 PROTEIN/DNA) FIVE. [1]	לחון כיווים יהיהי יחים יוים אינוס אינוס
Compound		YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS P5	INITIATOR ELEMENT	DNA; CHAIN: A, B;				÷	ADRI; CHAIN: NULL;					ADR1; CHAIN: NULL;				ADRI; CHAIN: NULL;			•		ADRI; CHAIN: NULL;				ADRI: CHAIN: NIII I	Mari, Cimer, Notae,	×.			ZINC FINGER PROTEIN GLII; CHAIN: A; DNA;	CITALLY: C, L,
SEQ FOLD score	·								51.79					÷								-				. •							
PMF score		1.00										,		0.25				60.0					0.09				-0.01	100			•	-0.15	
Verify score		0.22												-0.35				-0.30					0.41				0.10	}				-0.00	
Psi Blast		1.7e-34							6.8e-15					6.8e-15				8.5e-08					3.4e-14		٠		1 76-17	71.	-			3.4e-21	
AA END		200							506					231				06					341				356	2				355	
START AA		88							148					176	-			61					278				315	3				183	1
CHAIN		ပ																														¥	
PDB ID		lubd							2adr					2adr				2adr					2adr				2adr					2gli	
SEQ ID NO:	•	306							306					306				306					306				306					306	

Table

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PDB annotation	FINGER, COMPLEX (DNA-BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA-FINGER)	BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-	FINGER GLI; GLI, ZINC	BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEINE) FIVE.	FINGER GEL; GLI, ZINC	FINGER, COMPLEX (DNA-BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING	PROTEIN/DNA) FIVE-	FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA-	BINDING PROTEIN/DNA)		349	IMMUNOGLOBULIN, FAB.	ANTIBODY, ANTI-B-	COMPLEX (VIRAL	CAPSID/IMMUNOGLOBULIN	P24; FAB, FAB LIGHT	CHAIN, FAB HEAVY CHAIN	CAPSID/IMMUNOGLOBULIN
Compound		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA;	CHAIN: C, D;		ZINC FINGER PROTEIN	CHAIN: C, D;		ZINC FINGER PROTEIN	GLII; CHAIN: A; DNA;	CHAIN: C, D;		ELECTRON TRANSFER(IRON-SULFUR PROTEIN) RUBREDOXIN GRYN 3	C LTONIO	MONOCLONAL ANTI-E- SELECTIN 7A9	ANTIBODY; CHAIN: L, H;	HUMAN	IMMUNODEFICIENCY VIRIS TYPE I CAPSID	CHAIN: A, B; ANTIBODY	FAB25.3 FRAGMENT;	(17) To 17) To 14)
SEQ FOLD score	·			101.31												50.83					-	
PMF score		0.48					00'1			0.72				96:0				96.0				
Verify score	-	0.19					0.32			0.12				0.63				-0.14				
Psi Blast	·	1.7e-26		1.2e-63			1.2e-63			1.7e-32				0.0036		5.1c-18		le-18				
END A:A		143		202			200			228				379		216		207				
START		26		63			92			86				346		2		25				
CHAIN		ď		∢			Ą			Ą						н		H		٠.	,	
PDB U		2gli		2gli	-		2gli			2gli				6rxn		la5f		lafv				
SEQ ID NO:		306		306			306			306				306		307		307				

Table

				_								_							_		-,- -				_			_
PDB annotation	P24	COMPLEX (MHC/VIRAL PEPTIDE/RECEPTOR) HLA-A2 HRAVY CHAIN: CLASS I	MHC, T-CELL RECEPTOR,	COMPLEX MHCVVRAL	PEPTIDE/RECEPTOR	IMMUNOGLOBULIN	TETANUS TOXOID, HIGH	AFFINITY, CRYSTAL 2	PACKING MOTIF,	PROGRAMMING PROPENSITY TO	CRYSTALLIZE, 3	IMMUNOGLOBULIN	IMMUNOGLOBULIN.	ANTIBODY, FAB, ENZYME	START	À					CATALYTIC ANTIBODY	CATALYTIC ANTIBODY, [1]	TERPENOID SYNTHASE, **	CARBOCATION, 2	CYCLIZATION CASCADE	IMMUNE SYSTEM ANTI-	FAB 3F4 ANTI-PRION	ANTIBODY, FAB 3F4
Compound		HLA-A 0201; CHAIN: A; BETA-2 MICROGLOBULIN; CHAIN: B: TAX PEPTIDE.	CHAIN: C; T CELL	CHAIN: D. T. CHI.	RECEPTOR BETA; CHAIN: E;	FAB B7-15A2; CHAIN: L, H;		•					TP7 FAB; CHAIN: L, H;			COMPLEX	(ANTIBODY/ANTIGEN) HYHEL-5 FAB	COMPLEXED WITH	BOBWHITE QUAIL	LYSOZYME IBQL 3 IBQL	CATALYTIC ANTIBODY	19A4 (LIGHT CHAIN);	CHAIN: L; CATALYTIC	ANTIBODY 19A4 (HEAVY	CHAIN); CHAIN: H;	FAB ANTIBODY LIGHT	ANTIBODY HRAVY	CHAIN; CHAIN: H;
SEQ FOLD score		52.66				51.01				•			53.04								51.50							
PMF score								,								0.18										0:30		
Verify				•												0.00										0.10		
Psi Blast		0.0036				8.5e-16							1.5e-15			5.1e-18					3.4e-17	}	• • •			1.5e-19		
END AA		214				210							216			207				_	218)	٠			211		
START AA		22				13							5			33					5					31		
CHAIN		凹				J							н			Ħ					Н	!				H		
PDB U		lao7		_		laqk		-					layl			1bq1					1cf8	}				1cr9		
SEQ ID NO:		307				307							307		,	307					307	3				307		

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PDB annotation	CELL ADHESION NEURAL CELL ADHESION	GROWTH FACTOR/GROWTH PACTOR RECEPTOR FGF, FGFR, IMMUNOGLOBULIN- LIKE, SIGNAL	TRANSDUCTION, 2 DIMERIZATION, GROWTH FACTOR/GROWTH FACTOR RECEPTOR			VIRUS/VIRAL PROTEIN,	HUMAN POLIOVIRUS,	POLIOVRUS-RECEPTOR COMPLEX, VIRUS/VIRAL	PROTEIN, RECEPTOR	IMMUNE SYSTEM ANTI- LYSOZYME ANTIBODY,	HYHEL-63, HEN EGG WHIT		· ·	IMMUNE SYSTEM ANTI- FULLERENE ANTIBODY.	NANOTUBES	IMMUNE SYSTEM ANTI-	SWEETENER FAB, ANTIGEN-ANTIBODY,
Compound	AXONIN-1; CHAIN: A;	FIBROBLAST GROWTH FACTOR 2; CHAIN: A, B; FIBROBLAST GROWTH FACTOR RECEPTOR 1;	CHAIN: C, D;	IMMUNOGLOBULIN FAB' FRAGMENT OF THE DB3 ANTI-STEROID MONOCLONAL	ANTIBODY IDBB 3 (IGG1, SUBGROUP 2A, KAPPA 1) COMPLEX WITH PROFESTER OWE 1DBB 4.	POLIOVIRUS RECEPTOR;	VP2; CHAIN: 2; VP3; CHAIN: 3: VP4: CHAIN: 4:	(T. ', ', ', ', ', ', ', ', ', ', ', ', ',		ANTI-LYSOZYME ANTIBODY HYHEL-63	(LIGHT CHAIN); CHAIN: A,	ANTIBODY HYHEL-63	(HEAVY CHAIN); CHAIN: B. D;	IGG ANTIBODY (LIGHT CHAIN): CHAIN: L. IGG	ANTIBODY (HEAVY CHAIN); CHAIN: H;	FAB NC10.14 - LIGHT	CHAIN; CHAIN: L, A; FAB NC10.14 - HEAVY CHAIN;
SEQ FOLD score							٠.										
PMF score	0.48	0.09		0.05		0.31				0.24				0.83		69:0	
Verify score	-0.13	0.01		-0.40		-0.45				-0.02				0.19		90:0	
Psi Blast	3.4e-23	5.1e-17		5.1e-19		1.7e-25				5.1e-19				3.4e-19		1.4e-16	
END	211	209		207		218				207				207		212	
START AA	76	37		25		23				25				25		36	
CHAIN	Ą	ບ		H	-	R				—				H		H	
EDB EDB	1cs6.	Icvs		1dbb		1dgi				1 d qq		•		lemt		letz	
SEQ ID NO:	307	307		307		307				307				307		307	

				<u>·</u>										
PDB annotation	COMPLEX, CRYSTAL 2 STRUCTURE, RECEPTOR MIMICRY, ANTIGEN RECOGNITION	GROWTH FACTOR/GROWTH FACTOR RECEPTOR FGF2; FGFR2; IMMUNOGLOBULIN (IG)LIKE DOMAINS BELONGING TO THE L-SET 2	SUBGROUP WITHIN IG-LIKE DOMAINS, B-TREFOIL FOLD GROWTH FACTOR/GROWTH	FACTOR RECEPTOR FGF1; FGFR1; IMMUNOGLOBULIN (IG) LIKE DOMAINS BELONGING TO THE I-SET 2 STIRGEOTE WITHIN IG-1 IKP	DOMAINS, B-TREFOIL FOLD	IMMUNOGLOBULIN,	ANTIBODY, FAB, HEPATITIS B, PRES2		CATALYTIC ANTIBODY, AMIDINIUM, HAPTENIC CHARGE	IMMUNE SYSTEM IMMUNOGLOBULIN FOLD (f)	02/		IMMUNOGLOBULIN H	IMMUNOGLOBULIN ÎŢ
Compound	CHAIN: H, B	FIBROBLAST GROWTH FACTOR 2; CHAIN: A, B, C, D; FIBROBLAST GROWTH FACTOR RECEPTOR 2; CHAIN: E, F, G, H:	FIBROBLAST GROWTH	FACTOR 1; CHAIN: A, B; FIBROBLAST GROWTH FACTOR RECEPTOR 1; CHAIN: C, D;		F124 IMMUNOGLOBULIN (KAPPA LIGHT CHAIN);	CHAIN: A, C; F124 IMMUNOGLOBULIN (IGG1 HEAVY CHAIN); CHAIN: B, D:	CATALYTIC ANTIBODY	482; CHAIN: L, J; CATALYTIC ANTIBODY 482; CHAIN: H, K;	BLUE FLUORESCENT ANTIBODY (19G2)-HEAVY	CHAIN; CHAIN: H, A; BLUE FLUORESCENT ANTIBODY (1902)-LIGHT	CHAIN; CHAIN: L, B;	4-4-20 (IG*G2A=KAPPA=) FAB FRAGMENT; IFLR 5 CHAIN: L, H; 1FLR 6	ANTIBODY M41; CHAIN: L, H, M, I;
SEQ FOLD score														
PMF score		0.03	0.05			0.90		0.57		0.65	,		0.42	0.57
Verify score		-0.02	0.08			0.20		0.11		-0.12			-0.04	0.07
Psi Blast		5.1e-17	1.2e-17			1e-18		1.7e-18		6.8e-20			1.7e-18	3.4e-20
END AA		199	209			202		207		207			207	207
START AA		37	37			22	<u>-</u>	25		31			31	25
CHAIN		Ð	ပ			æ		H		н			Ħ	H
PDB ED		lev2	levt			1211		1£3d		1fl3			41	Igpo
SEQ ID NO:		307	307			307		307		307			307	307

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PDB annotation	ANTIBODY DESIGN, IMMUNOGLOBULIN 2 STRUCTURE, ANTIGEN- BINDING SITE, CANONICAL CONFORMATION, 3 COMPLEMENTARITY- DETHRMINING REGION:	IMMUNOGLOBULIN INTACT IMMUNOGLOBULIN, V REGION, C REGION, HINGE REGION			COMPLEX (IMMUNOGLOBULINIHYDR) OLASE) N10 FAB IMMUNOGLOBULIN; INSN [] STAPHYLOCOCCAL NUCLEASE NUCLEASE RIBONUCLEATE, INSN 11 IMMUNOGLOBULIN, STAPHYLOCOCCAL NUCLEASE INSN 25	MONOCLONAL ANTIBODY F
Compound		IGGI INTACT ANTIBODY MAB61.1.3; CHAIN: A, B, C, D	IMMUNOGLOBULIN CHA255 IMMUNOGLOBULIN FAB' FRAGMENT (IGGI- LAMBDA) COMPLEX 1IND 3 WITH 4-[N-(2- HYDROXYETHYL)- THIOUREIDO]-L-BENZYL- 1IND 4 EDTA-IN(3+) (INDIUM(3+)-EOTUBE)	IMMUNOGLOBULIN FAB D44.1 (IGG1,KAPPA) (BALB/C MOUSE, MONOCLONAL ANTIBODY) 1MLB 5	IGG FAB (IGG1, KAPPA); INSN 4 CHAIN: L, H: INSN 5 STAPHYLOCOCCAL NUCLEASE; INSN 9 CHAÎN: S; INSN 10	MONOCLONAL ANTIBODY 3A2; CHAÎN: H,
SEQ FOLD score	-					
PMF		0.43	0.51	0.45	0.84	0.54
Verify score		-0.29	0.09	0.31	0.03	-0.07
Psi Blast		3.4e-18	1.46-18	3.4e-17	5.1e-19	1.7e-19
AA AA		207	207	207	207	207
START AA		31		33	22	31
CHAIN		В	Ħ	В	н	н
PDB ID		ligy	lind	Imlb	Insn	1sbs
SEQ ID NO:		307	307	307	307	307

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PDB annotation	FAB-FRAGMENT, REPRODUCTION	COMPLEX (ANTIBODY/PEPTIDE	EFITOFE) ANTIGEN, PEPTIDE ANTIGEN,	ANTITUMOR ANTIBODY, 2	COMPLEX (ANTIBODY/PEPTIDE EPITOPE)	COMPLEX	(ANTIBODY/PEPTIDE EPITOPE) ANTIBODY.	PEPTIDE ANTIGEN,	ANTITUMOR ANTIBODY, 2	COMPLEX (ANTIBODY/PEPTIDE	EPITOPE)					CATALYTIC ANTIBODY [7]	CATALYTIC ANTIBODY,	REACTION				a veri			and the second	
Compound	L;	SM3 ANTIBODY; CHAIN: L, H; PEPTIDB EPITOPE;	CHAIN: F;			SM3 ANTIBODY; CHAIN:	L, H; PEPTIDE EPITOPE; CHAIN: P:					IMMUNOGLOBULIN IGGI	FRAGMENT (TE33)	COMPLEX WITH	PEPTIDE 3 (CTP3) 1TET 4	IGG 5C8; CHAIN: L, H;			IMMUNOGLOBULIN	ED A GIVENT COMPI EXED	WITH ANTIGEN 2CGR 3 N-	(P-CYANOPHENYL)-N-	(DIPHENYLEMETHYL)	GUANIDINEACETIC ACID 2CGR 4	COMPLEX(ANTIBODY-	ANTIGEN) IG*G1 FAB FRAGMENT (HY/HEL\$-10)
SEQ FOLD score	-			,		53.74						-								-						
PMF score		0.94		. •								0.30				0.82			0.18						0.39	
Verify score		-0.03		٠								-0.32				-0.18			0.28						-0.16	
Psi Blast		1e-20				5.1e-15	-			· .		3.46-19				1.7e-18			1.7e-16	-					1e-20	-
END		207				206						207				207			207						207	
START		31				22						22				31			33						25	•
CHAIN		н				I	,			_		н				Ħ			H						H	-
PDB CD		1sm3				1sm3						Itet				25c8			2cgr						3hfm	
SEQ ID		307				307						307				307			307				-		307	

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PDB annotation			SIGNALING PROTEIN GTP- BINDING PROTEINS, PROTEIN-PROTEIN	COMPLEX, EFFECTORS	SIGNALING PROTEIN GTP- BINDING PROTEINS,	PROTEIN-PROTEIN COMPLEX, EFFECTORS		SIGNALING PROTEIN GTP-	BINDING PROTEIN RHOA,	GIFASE KHOA; KHO GDI 1; RHO GTPASE G-PROTEIN.	SIGNALING PROTEIN	SIGNALING PROTEIN G	٠.	CBVSTAIL OGB ABHY 2	SIGNALING PROTEIN	SIGNALING PROTEIN G	PROTEIN, GTP	HYDROLYSIS, KINETIC	SIGNAL ING PROTEIN	SIGNALING PROTEIN IL	PROTEIN-PROTEIN	COMPLEX, ANTIPARALLEI	S/EXOCYTOS	ASE,	RAB6, VESICULAR TRAFFICKING FIL
Compound	AND LYSOZYME (B.C.3.2.1.17) 3HFM 4 COMPLEX 3HFM 5		RAS-RELATED PROTEIN RAP-1A; CHAIN: A; PROTO-ONKOGENE	SERINE/THREONINE PROTEIN KINASE CHAIN: B;	RAS-RELATED PROTEIN RAP-1A; CHAIN: A;	PROTO-ONKOGENE SERINE/THREONINE	PROTEIN KINASE CHAIN: B;	TRANSFORMING PROTEIN	RHOA; CHAIN: A, C; RHO	INHIBITOR ALPHA:	CHAIN: E, F;	TRANSFORMING PROTEIN	P21/H-RAS-1; CHAIN: A;			TRANSFORMING PROTEIN	P21/H-RAS-1; CHAIN: A;		-	HIS-TAGGED	TRANSFORMING PROTEIN	KHOA(0-181); CHAIN: A; PKN: CHAIN: B:	RAB6 GTPASE; CHAIN: A;		
SEQ FOLD score			57.43				,		•			51.73													
PMF score	·				0.45			0.00								0.36				0.16			0.25		
Verify score					0.28			0.04								0.34				0.34			0:30		
Psi Blast			1.2e-63		1.2e-63			1.2e-51				1e-65				1e-65				5.1e-53			3.4e-54	•	
END A			163	•	163			171				164				163				163			161		
START AA			I		3			4				1				3				4			3		
CHAIN	·		¥		V			¥				Ą				¥				A			Ą		
PDB ID			lcly		lcly			1cc0				lctq				lctq				1cxz			1d5c		·
SEQ ID NO:			309		309			309				309				309				309			309	-	·

Table 5

PDB annotation		TRAFFIC, GTP HVDBOI VSIS VPT/RAB 2	PROTEIN, ENDOCYTOSIS,	HYDROLASE	GTP-BINDING PROTEIN	GIP-BINDING PROTEIN,	GDP, RAS	GTP-BINDING PROTEIN	GTP-BINDING PROTEIN,	SMALL G PROTEIN, RAP2, GDP, RAS	GTP-BINDING GTP-	BINDING, GTPASE, SMALL	G-PROTEIN, RHO FAMILY, RAS SUPER 2 FAMILY	-		KELATED PROTEIN KABSA;	BINDING/FEFFOR) G	PROTRIN EFFECTOR	<i></i>	EXOCYTOSIS, RAB	PROTEIN, RAB3A,	KABPHILIN	HYDROLASE G PROTEIN, (F)	GTP HYDROLYSIS, RAB 2	PROTEIN,	NEUROTRANSMITTER	RELEASE, HYDROLASE	222	IN; PEPTIDE/RECEPTOR) HLA- [1] A2 HEAVY CHAIN; CLASS I[1]
Compound	GTP-BINDING PROTEIN YPT51; CHAIN: A;		-		RAP2A; CHAIN: NULL;		-	RAP2A; CHAIN: NULL;			RACI; CHAIN: NULL;			RAB-3A; CHAIN: A;	RABPHILIN-3A; CHAIN: B;			•		•			RAB3A; CHAIN: A;		, •		-		HLA-A 0201; CHAIN: A; BETA-2 MCROGLOBULN; CHAIN: B; TAX PEPTIDE;
SEQ FOLD score					68.65							. 1								٠			•			-			73.76
PMF score	0.29							0.72			-0.01	•		0.34	-				,				0.36						
Verify score	0.41			,				0.14			0.10			0.05		-							-0.08		-				
Psi Blast	3.4e-51				6.8e-60			6.8e-60			3.4e-52			6.8e-61				-					1.7e-61						2.4e-32
END AA	164				164			160			166			167						á			<u>2</u>						118
START AA	2				1			3			3		•	3			,						m						23
CHAIN	¥													Ą									∢						Q
PDB UD	1ek0				1kao			Ikao	_		1mh1			1zbd									3rab			-			1807
SEQ ID NO:	309				309			309			309			309									309						310

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PDB annotation	VIRAL PEPTIDE, 2 COMPLEX (MHC/VIRAL PEPTIDE/RECEPTOR	IMMUNE SYSTEM BENCE- JONES; IMMUNOGLOBULIN, AMYLOID, IMMUNE SYSTEM	IMMUNOGLOBULIN IMMUNOGLOBULIN, KAPPA LIGHT-CHAIN DIMER HEADER	T CELL RECEPTOR TCR; T CELL RECEPTOR, MHC CLASS I, HUMAN IMMUNODEFICIENCY VIRUS, 2 MOLECULAR RECOGNITION	COMPLEX (MHC/VIRAL PEPTIDE/RECEPTOR) HLA A2 HEAVY CHAIN; COMPLEX (MHC/VIRAL PEPTIDE/RECEPTOR)	COMPLEX (ANTIBODY/ANTIGEN) FAB. 12; VEGF, COMPLEX (ANTIBODY/ANTIGEN), (I) ANGIOGENIC FACTOR	IMMUNE SYSTEM REIV, TUSTABILIZED IMMUNOGLOBULIN FRAGMENT, BENCE-JONES 2 PROTEIN, IMMUNE SYSTEM	IMMUNE SYSTEM REIV, IU STABILIZED
Compound	RECEPTOR ALPHA; CHAIN: D; T CELL RECEPTOR BETA; CHAIN: E;	BENCE-JONES KAPPA I PROTEIN BRE; CHAIN: A, B, C;	IMMUNOGLOBULIN; CHAIN: A, B;	T CELL RECEPTOR V. ALPHA DOMAIN; CHAIN: A, B;	HLA-A 0201; CHAIN: A; BETA-2 MICROGLOBULIN; CHAIN: B; TAX PEPTIDE; CHAIN: C; T CELL RECEPTOR ALPHA; CHAIN: D; T CELL RECEPTOR BETA; CHAIN: B;	FAB FRAGMENT; CHAIN: L, H, J, K; VASCULAR ENDOTHELIAL GROWTH FACTOR; CHAIN: V, W;	IG KAPPA CHAIN V-I REGION REI; CHAIN: A, B;	IG KAPPA CHAIN V-I REGION REI; CHAIN: A, B;
SEQ FOLD score		57.66		87.03	55.59		54.30	
PMF score			0.95			0.99		0.99
Verify score			0.21			0.24		0.51
Psi Blast		5.ie-43	1.7e-44	8.5e-40	1e-34	3.4e-46	1.2e-43	1.2e-43
END AA		118	113	118	118	115	118	113
START AA	·	23	24	23	23	42	20	24
CHAIN		Ą	Ą	¥	Q	٦	∀	A
PDB ID		150w	1b6d	1b88	1bd2	16j1	1bww	1bww
SEQ ID NO:		310	310	310	310	310	310	310

Table 5

PDB annotation	IMMUNOGLOBULIN FRAGMENT, BENCE-JONES 2 PROTEIN, IMMUNE SYSTEM	IMMUNE SYSTEM FAB-IBP COMPLEX CRYSTAL STRUCTURE 2.7A RESOLUTION BINDING 2 OUTSIDE THE ANTIGEN COMBINING SITE SUPERANTIGEN FAB VH3 3 SPECIFICITY		IMMUNE SYSTEM IMMUNOGLOBULIN FOLD, ANTIBODY, IGM, FV	Ď	T 450	2/0.	
Compound		IGM RF 2A2; CHAIN: A, C, E; IGM RF 2A2; CHAIN: B, D, F; IMMUNOGLOBULIN G BINDING PROTEIN A; CHAIN: G, H;	IMMUNOGLOBULIN 3D6 PAB 1DFB 3	IGM MEZ IMMUNOGLOBULIN; CHAIN: L; IGM MEZ IMMUNOGLOBULIN; CHAIN: H;	IMMUNOGLOBULIN FV FRAGMENT OF A HUMANIZED VERSION OF THE ANTI-CD18 1FGV 3 ANTIBODY FIS2' (HUH52- AA FV) 1FGV 4	IMMUNOGLOBULIN FV FRAGMENT OF A HUMANIZED VERSION OF THE ANTI-CD18 IFGV 3 ANTIBODY H52' (HUH52- AA FV) 1FGV 4	IMMUNOGLOBULIN IMMUNOGLOBULIN M (IG-M) FV FRAGMENT IIGM 3	IMMUNOGLOBULIN IMMUNOGLOBULIN M (IG-M) FV FRAGMENT IIGM 3
SEQ FOLD score					55.38		52.10	·
PMF score		0.98	1.00	66:0		0.99		1.00
Verify score		0.12	0.43	0.66		0.37		0.42
Psi Blast		1e-47	3.4e-46	5.1e-44	1.7e-46	1.7e-46	1.7e.44	1.7e-44
END		113	113	113	117	113	118	113
START AA		24	74	24	23	24	23	24
CHAIN		¥	L	I	ı	ļ i	1	1
PDB ID		1dee	1dfb	lqbl	1fgv	lfgv	ligm	ligm
SEQ ID NO:		310	310	310	310	310	310	310

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PDB annotation	COMPLEX (IMMUNOGLOBULINIRECEP. TOR) TCR VAPLHA VBETA DOMAIN; T-CELL. RECEPTOR, STRAND SWITCH, FAB, ANTICLONOTYPIC, 2 (IMMUNOGLOBULINIRECEP. TOR)	IMMUNOGLOBULIN TR1.9, ANTI-THYROID PEROXIDASE, AUTOANTIBODY, 2 IMMUNOGLOBULIN			TRANSFERASE TRANSFERASE, HERBICID起 DETOXIFICATION	COMPLEX (TRANSFERASE/LIGAND) (COMPLEX (TRANSFERASE/LIGAND), COMPLEX (TRANSFERASE, HERBICIDE) 2 DETOXIFICATION HEADER	RINCLEOTIDE
Compound	KBS-C20 T-CELL ANTIGEN RECEPTOR; CHAIN: A, B; ANTIBODY DESIRB-1; CHAIN: L, H;	TRI.9 FAB; CHAIN: L, H;	IMMUNOGLOBULIN WAT, A VARIABLE DOMAIN FROM IMMUNOGLOBULIN LIGHT-CHAIN 1WTL 3 (BENCE-JONES PROTEIN) 1WTL 4	IMMUNOGLOBULIN FAB FRAGMENT OF A HUMANIZED VERSION OF THE ANTI-CD18 2FGW 3 ANTIBODY H52' (HUH52- OZ FAB) 2FGW 4	GLUTATHIONE S- TRANSFERASE III; CHAIN: NULL;	GLUTATHONE S- TRANSFERASE I; CHAIN: A, B; LACTOYLGLUTATHONE; CHAIN: C, D	ELONGATION FACTOR 1-
SEQ FOLD score	84.21		50.36				137.96
PMF score		96'0		0.93	0.04	-0.09	
Verify		0.55		0.43	0.01	0.17	
Psi Blast	5.1e-42	1.5e-43	1.7e-42	3.4e-46	5.1e-14	1.4e-14	1.4e-46
END	118	113	118	113	78	89	225
START AA	23	25	23	24	1		135
CHAIN	4	1	Ą	1		V	
PDB U	1,465	lvge	lwti	2fgw	law9	laxd	1b64
SEQ ID NO:	310	310	310	310	311	311	311

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PDB annotation	EXCHANGE FACTOR GUANINE NUCLEOTIDE EXCHANGE FACTOR, G- PROTEIN, TRANSLATION 2	ELONGATION	GUANINE NUCLEOTIDE EXCHANGE FACTOR	GUANINE NUCLEOTIDE	EXCHANGE FACTOR, G-	PROTEIN, TRANSLATION 2	SIGNAL PROTEIN NUCLEAR	SIGNAL PROTEIN	ANKYRIN BINDING MAB;	ANKYKIN BINDING,	TRANSFERASE, CARRIER 2	CRYSTALLIZATION, ION	TRANSPORT	TRANSFERASE, BLOOD	CLOTTING GAMMA CHAIN	CAPPED PROTEIN DEIVEN	2 CRYSTALLIZATION	TRANSLATION PROTEIN-	PROTEIN COMPLEX		GST;	TRANSFERASE,	DETOXIFICATION, (IL)	GLUTATHIONE TRANSFERASE			Trace	il.
Compound	BETA; CHAIN: NULL;		ELONGATION FACTOR 1- BETA; CHAIN: NULL;			:	AML-1B; CHAIN: A;		FUSION PROTEIN OF	ALPHA-NA,K-AIPASE	WILD CRAIN: INOLL;			CHIMERA OF	GLUTATHIONE S-	SYNTHETIC CHAIN: A B.	SINIMBIIC CHAIN: A, B;	ELONGATION FACTOR	EEF1A; CHAIN: A;	ELONGATION FACTOR FEFIBA: CHAIN: B:	GLUTATHIONE	TRANSFERASE; CHAIN:	NULL;		GLUTATHIONE	TRANSFERASE	TRANSFERASE	(E.C.2.3.1.16) FUSED WITH
SEQ FOLD score																												
PMF			8: 				0.27	•	0.33					0.27				1.00			0.31				0.12			
Verify			0.43				0.24		0.28					0.19				0.20			0.07				0.26			
Psi Blast		,	1.46-40				5.1e-13	•	5.1e-13					5.1e-13			i	4.8e-48			3.4e-16			•	5.10-13			
END AA		900	C77				91		91					91			-	225			22				16		•	
START			142				5		5					S				137			2				5			
CHAIN				,			Ą							⋖			İ	В										
PDB ID		1771	1004				1b8x		1bg5	-				ldug				1f60			1fhe	. — 	- 		Igne			
SEQ ID NO:			311				311		311					311				311			311		,		311			

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				PETSUSOE	/01	
PDB annotation			TRANSFERASE TRANSFERASE, GLUTATHIONE, CONJUGATION, DETOXIFICATION, CYTOSOLIC, DIMER	GENE REGULATION POZ DOMAIN; PROTEIN- PROTEIN INTERACTION DOMAIN, TRANSCRIPTIONAL 2 REPRESSOR, ZINC-FINGER PROTEIN, X-RAY CRYSTALLOGRAPHY, 3 PROTEIN STRUCTURE, PROMYELOCYTIC LEUKEMIA, GENE REGULATION		COMPLEX (ZINC FINGER/DNA) COMPLEX
Compound	A IGNE 3 CONSERVED NEUTRALIZING EPITOPE ON GP41 OF HUMAN IGNE 4 IMMUNODEFICIENCY VIRUS TYPE 1, COMPLEXED WITH GLUTATHIONE IGNE 5	GLUTATHIONE TRANSFERASE GLUTATHIONE S- TRANSFERASE (E.C.2.5.1.18) (26 KDA) 1GTA 3	GLUTATHIONE S- TRANSFERASE; CHAÎN: A, B, C, D;	PROMYELOCYTIC LEUKEMIA ZINC FINGER PROTEIN PLZF; CHAIN: A;	OXIDOREDUCTASB(OXYG EN(A)) GALACTOSE OXIDASE (B.C.1.1.3.9) (PH 4.5) 1GOF 3	QGSR ZINC FINGER PEPTIDE; CHAIN: A;
SEQ FOLD score		·			·	
PMF score		0.24	-0.01	86.0	0.01	0.36
Verify		-0.04	0.15	0.19	0.25	-0.01
Psi Blast		5.1e-13	5.1e-15	1.4e-25	1.7e-12	2.4e-15
END AA		91	986	164	615	220
START		رح د	ъ	40	365	118
CHAIN			∀	4	-	A
PDB CD		1gta	1gtu	1buo	lgof	laih
SEQ ID NO:	·	311	311	312	312	313

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PDB annotation	(ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING FINGER, DNA-BINDING FINGER, DNA-BINDING	COMPLEX (ZINC FINGER/DNA) COMPLEX (I (ZINC FINGER/DNA), ZINC (I FINGER, DNA-BINDING (I PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX FINGER/DNA) COMPLEX FINGER/DNA), ZINC [L] FINGER, DNA-BINDING [L] PROTEIN
Compound	DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B,	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	OGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B,
SEQ FOLD score				:			
PMF score		-0.13	0.17	0.53	0.74	1.00	1.00
Verify		0.03	-0.26	-0.03	-0.02	0.17	0.28
Psi Blast		3.46-24	5.1e-28	8.5e-24	8.4e-30	66-35	6.8e-31
END AA		219	247	303	332	329	387
START AA		144	167	561	256	279	307
CHAIN		¥	¥	V	¥	¥	¥
PDB ID		lalh	laih	lalh	lalh	lalh	laih
SEQ ID NO:		313	313	313	313	313	313

Table :

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PDB annotation		COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN		COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN (I) DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER PROTEIN-DNA) ZINC FINGER PROTEIN-DNA INTERACTION, PROTEIN IN DESIGN, 2 CRYSTAL IN STRUCTURE, COMPLEX IN
Compound	Ċ	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	DNA-BINDING PROTEIN HUMAN ENHANCER- BINDING PROTEIN MBP-1 MUTANT WITH CYS 11 1BBO 3 REPLACED BY ABU (C11ABU) (NMR, 60 STRUCTURES) 1BBO 4	DNA; CHAIN: A, B, D, E, CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E, CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;
SEQ FOLD score				76.59			
PMF score		0.99	0.87		0.06	0.15	0.52
Verify		0.35	-0.15		0.01	0.18	-0.23
Psi Blast		1.2e-39	6e-39	6e-39	1.1e-15	1.7e-41	3.46-47
END AA		388	472	473	307	219	247
START		307	363	391	254	143	166
CHAIN		A	A	A		υ	υ
PDB TD		laih	laih	laih	1bbo	1теу	1mey
SEQ ID NO:		313	313	313	313	313	313

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PDB annotation	(ZINC FINGER/DNA)	COMPLEX (ZINC	PROTEIN-DNA	INTERACTION, PROTEIN	DESIGN, 2 CRYSTAL	STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC	FINGER/DNA) ZINC FINGER,	PROTEIN-DNA	INTERACTION, PROTEIN	DESIGN, 2 CRYSTAL	STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC	FINGER/DNA) ZINC FINGER,	PROTEIN-DNA	INTERACTION, PROTEIN	DESIGN, 2 CRYSTAL	STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC	FINGER/DNA) ZINC FINGER,	PROTEIN-DNA	INTERACTION, PROTEIN	DESIGN, 2 CRYSTAL	STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC	FINGER/DNA) ZINC FINGER,	FROI ELIVEDIA DE CITETAL	INTERACTION, PROTEIN A	DESIGN, 2 CRISIAL	STRUCTURE, COMPLEX	CONTROL BY CONTROL	FINGER/DNA) ZINC FINGER, F	FROI BIN-DIA
Compound		DNA; CHAIN: A, B, D, E;	PROTEIN; CHAIN: C, F, G;					DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G,					DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;					DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;	•		•		DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER PROTEIN: CHANGE	INCIDENT, CHAMIN. C. F. G.				DNA: CHAIN: A B D E.	CONSENSUS ZINC FINGER	FROIDIN; CHAIN: C, F, G;
SEQ FOLD score										,					·	•																	-				
PMF score		0.46						1.00		,			-		1.00							1.00							90.						5	3	
Verify score		0.27						0.31							0.20							0.22							0.24		•				030	3	
Psi Blast		6.8e-45	,					1.7e-50		,	-				3.4e-51						,	3.4e-51							8.5e-51						5 10.51		7
END		331			,			359							387							415							443						471	:	7
START AA		222						278							306	,						334							362						390		
CHAIN		၁						ပ							ບ							၁							ပ						ر)	
PDB ED		1mey						Imey							1mey							lmey							Imey						1mev		
SEQ ID NO:		313						313	•						313							313							313.			•		•	313	}	

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PDB annotation	INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DINA) ZINC FINGER	PROTEIN-DNA	INTERACTION, PROTEIN	DESIGN, 2 CRYSTAL	STRUCTURE, COMPLEX	COMPLEX (ZINC	FINGER/DNA) ZINC FINGER,	PROTEIN-DNA	INTERACTION, PROTEIN	DESIGN, 2 CRYSTAL	STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC	FINGER/DNA) ZINC FINGER,	INCIDENTAL ACTIONS	INTERACTION, PROTEIN	DESIGN, 2 CKISIAL	STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	PECTI ATTOMONANTION	SCOLATION/DINA) IFILIA;	PROTTEIN TINA	TRANSCRIPTION PACTOR.	5S RNA 2 GENE. DNA	BINDING PROTEIN, ZINC	FINGER, COMPLEX 3	(TRANSCRIPTION	REGULATION/DNA)	COMPLEX (TRANSCRIPTIONF REGULATION/DNA) TFIIIA; [1	5S GENE; NMR, TFIIIA,	PROTEIN, DNA,
Compound			DNA; CHAIN: A, B, D, E;	PROTEIN; CHAIN: C, F, G;				DNA: CHAIN: A. B. D. E.	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;					DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER	TACLEMY, CARAMY, C, F, C,					IKANSCKIPTION FACTOR	GENE: CHAIN: A; 33 KINA GENE: CHAIN: H H:	Carrie Carrier Ly 13	•	-		•			TRANSCRIPTION FACTOR IIIA; CHAIN: A; 5S RNA	GENE; CHAIN: E, F;	
SEQ FOLD score			90.72																														
PMF score			-	•				1.00				-			96.0						1,7	C.43									0.04		
Verify score								0.09							0.03	•					,,,	-0.31									-0.17		
Psi Blast		;	5.le-51					3.4e-42	!						3.4e-13						1-16	CI-9I									7.2e-18		
END			472					484							331						200	S S									305		
START AA		3	390					418							304						2	3									196		
CHAIN		,	ပ					ပ							5							< -									≪		
PDB			Imey					Imey	`						Imey						871	CIIT									1		
SEQ ID NO:	•		313					313							313					-	212	CIC									313		

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PDB annotation	TRANSCRIPTION FACTOR, 5S RNA 2 GENE, DNA BINDING PROTEIN, ZINC FINGER, COMPLEX 3 (TRANSCRIPTION REGUL, ATTON/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) TFIIIA; 5S GENE; NMR, TFIIIA, PROTEIN, DNA, TRANSCRIPTION FACTOR, 5S RNA 2 GENE, DNA BINDING PROTEIN, ZINC FINGER, COMPLEX 3 (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INTIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION) REGULATION/DNA) COMPLEX (TRANSCRIPTION), REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INTITATION, ZINC FINGER FROTEIN	COMPLEX (TRANSCRIPTION: REGULATION/DNA) COMPLEX (TRANSCRIPTION) REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION TRANSCRIPTION
Compound		TRANSCRIPTION FACTOR IIIA; CHAIN: A; 5S RNA GENE; CHAIN: E, F;	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;
SEQ FOLD score			÷		÷
PMF		0.25	0.05	0.82	0.98
Verify		-0.04 40.04	-0.29	-0.11	0.11
Psi Blast		2.46-19	5.1e-31	8.5e-35	8.5e-38
END AA		333	312	966	431
START		258	167	268	279
CHAIN		Ą	V	V	∢
PDB D		द् <u>र</u> म	1466	1476	11f6
SEQ ID NO:		313	313	313	313

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PDB annotation	PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEA (I KANSCKIPIION REGULATION/DNA), RNA	POLYMERASE III, 2	TRANSCRIPTION	INITIATION, ZINC FINGER	PROTEIN	COMPLEX (TRANSCRIPTION PEGIT ATTONIONA)	CONDITION ON CENTRAL	REGIT ATTONONA). RNA	POLYMERASE III. 2	TRANSCRIPTION	INITIATION, ZINC FINGER	PROTEIN	COMPLEX (TRANSCRIPTION	REGULATION/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA), RNA	POLYMERASE III, 2	TRANSCRIPTION	INTITATION, ZINC FINGER	PROTEIN	COMPLEX (TRANSCRIPTION,	REGULATION/DNA) YING-	YANG 1; TRANSCRIPTION	INITIATION, INITIATOR	ELEMENT, YY1, ZINC 2	FINGER PROTEIN, DNA-	PROTEIN RECOGNITION, 3	COMPLEX (TRANSCRIPTION	REGULATION/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA) YING-	NETTATION INTERVED	ELEMENT, YYI, ZINC 2
Compound		TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE;	CERMIN: B, C, E, F;					TFIIIA; CHAIN: A, D; 5S	CHAIN: B C P P.	(1, t, t, t, t, t, t, t, t, t, t, t, t, t,			•	•	TFIIIA; CHAIN: A, D; 5S	RIBOSOMAL RNA GENE;	CHAIN: B, C, E, F;						YY1; CHAIN: C; ADENO-	ASSOCIATED VIRUS P5	INITIATOR ELEMENT	DNA; CHAIN: A, B;						YYI; CHAIN: C; ADENO-	ASSOCIATED VIRUS P5	DNA: CHAM: A B.	DIVA, CHAIN: A, B,
SEQ FOLD score		94.43									•						•		·																
PMF score								1.00		•			-		10.01	-	•						0.21									0.00			
Verify score							ļ	0.37							-0.37						-		-0.23									-0.17	•		
Psi Blast		1.7e-39					,	1.7e-39		•					8.5e-32								5.1e-29									3.4e-28			
END		485						485							228								247									303			
START AA		334					200	335							98								154									168			
CHAIN		¥						∢							٧								U									U			
PDB ID		1tf6						1110					•		1tf6				· · -				1ubd									Inbd			
SEQ ID NO:		313						313							313								313									313		•	

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PDB annotation	FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION	REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-	YANG 1; TRANSCRIPTION	ELEMENT, YYI, ZINC 2	FINGER PROTEIN, DNA-	PROTEIN RECOGNITION, 3	COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION	KEGULATION/DNA) YING- YANG 1: TRANSCRIPTION	INITIATION, INITIATOR	ELEMENT, YY1, ZINC 2	FINGER PROTEIN, DNA-	PROTEIN RECOGNITION, 3	REGILATION DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA) YING- "	YANG 1; TRANSCRIPTION	INITIATION, INITIATOR	ELEMENT, YY1, ZINC 2	FINGER PROTEIN, DNA-	PROTEIN RECOGNITION, 3	DEGIT ATTONONA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA) YING-	YANG 1; TRANSCRIPTION	INITIATION, INITIATOR	ELEMENT, YY1, ZINC 2	FINGER PROJEIN, DINA-	COMPLEX (TRANSCRIPTION)
Compound			YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS P5	INITIATOR ELEMENT	DNA; CHAIN: A, B;		:		YY1; CHAIN: C; ADENO-	ASSOCIATED VIRUS PS	DNA; CHAIN: A, B;					YY1; CHAIN: C; ADENO-	ASSOCIATED VIRUS P5	INITIATOR ELEMENT	DNA; CHAIN: A, B;					YY1; CHAIN: C; ADENO-	ASSOCIATED VIRUS P5	INTITATOR ELEMENT	DNA; CHAIN: A, B;			
SEQ FOLD score							-				·													79.23						
PMF			0.98						0.95							1.00						•						·		
Verify score			0.12						90.0							0.02								-						
Psi Blast			1.5e-33						1e-34		-					6.8e-35								3.4e-35						
END AA	·		329						387							415		٠.						472						
START AA			262						286			:				314					•			362						
CHAIN			ပ						ر ک							ပ								ပ						
PDB ID			lubd						pqnI							lubd				-				lubd						
SEQ ID NO:			313						313							313								313						

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PDB annotation	REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION	INITIATION, INITIATOR	FINGER PROTEIN, DNA-	PROTEIN RECOGNITION, 3	COMPLEX (TRANSCRIPTION REGIT ATTOMONA)	COMPLEX (TRANSCRIPTION	REGULATIONDNA) YING-	INITIATION, INITIATOR	ELEMENT, YY1, ZINC 2	FINGER PROTEIN, DNA-	PROTEIN RECOGNITION, 3	COMPLEX (IRANSCRIPTION DEGIT ATTOMONA)	COMPLEX (DNA-BINDING	PROTEIN/DNA) FIVE-	FINGER GLI; GLI, ZINC	FINGER, COMPLEX (DNA-	BINDING PROTRIN/DNA)	COMPLEX (DNA-BINDING	PROTEIN/DNA) FIVE	FINGER GLI; GLI, ZINC	FINGER, COMPLEX (DNA-	COMPLEX (DNA-BINDING	PROTEIN/DNA) FIVE-	FINGER GLI, GLI, ZINC	FINGER, COMPLEX (DNA-	BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING	FINGER GILI: GLI, ZINC	FINGER, COMPLEX (DNA-	BINDING PROTEIN/DINA)
Compound		YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT	DNA; CHAIN: A, B;	٠			YY1; CHAIN: C; ADENO-	ASSOCIATED VIRUS P5	DNA; CHAIN: A, B;					ZINC FINGER PROTEIN	GLII; CHAIN: A; DNA;	CHAIN: C, D;			ZINC FINGER PROTEIN	GLII; CHAIN: A; DNA;	CHAIN: C, D;		ZINC FINGER PROTEIN	GLII; CHAIN: A; DNA;	CHAIN: C, D;			ZINC FINGER PROTEIN	CHAIN: C. D.		
SEQ FOLD score		·			•																					•					
PMF score		1.00					1.00							0.43	!				0.05				0.50	3				1.00	,		
Verify		-0.19					-0.16							-0.16		-			-0.31				0.30	70.0				0.12			
Psi Blast		1e-34					5.1e-29		-					4.8e-29					5.1e-30				2 40.53	£:46-33				6e-54			
END		471					484							305					330				417	ì				389			
START		370					398							116					166				106	2				251			
CHAIN		ပ					ပ							▼					¥				<	۲				¥			
PDB		lubđ					lubd							2øli	6				2gli				:100	1187				2gli			
SEQ ID		313					313							313					313				212	crc				313			

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PDB annotation	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE- FINGER GLI, GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE- FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE- FINGER GLI, GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE- FINGER GI.I; GI.I, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEINDNA) FIVE- FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE- FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	KINASE RI(ALPHA); REGULATORY SUBUNIT, KINASE	TRANSCRIPTION/DNA COMPLEX (TRANSCRIPTION) REGULATION/DNA), DNA- BINDING, CAMP- 2
Compound	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	CAMP DEPENDENT PROTEIN KINASE; CHAIN: NUIL:	CATABOLITE GENE ACTIVATOR PROTEIN; CHAIN: A; DNA (5'- D(*GP*TP*CP*AP*CP*AP* TP*TP*AP*AP*T)-3');
SEQ FOLD score			88.39	·	·			
PMF score	0.98	0.98		0.98	66'0	0.87	0.81	0.35
Verify	0.19	0.22		-0.01	-0.10	-0.10	0.36	0.37
Psi Blast	8.5e-34	9.6e-64	9.66-64	3.4e-34	4.80-56	6.8e-30	2.6e-26	2.6e-22
END A	386	473	445	470	480	480	501	504
START	262	307	308	342	363	370	363	383
CHAIN	Ą	V	¥	Ą	¥	Ą		Ą
PDB UD	2gli	2gli	2gli	2gli	2gli	2gli	lrgs	2cgp
SEQ ID NO:	313	313	313	313	313	313	315	315

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PDB annotation		HYDROLASE SPASE I,	LEADER PEPTIDASE I;	SERINE-DEPENDANT	HYDROLASE. SIGNAL 2	PEPTIDE PROCESSING,	PROTEIN TRANSLOCATION,	MEMBRANE BOUND 3	PROTEIN ASE, MEMBRANE PROTEIN	HYDROLASE SPASE I,	LEADER PEPTIDASE I;	SERINE DEPENDANT	HYDROLASE, SIGNAL 2	PEPTIDE PROCESSING,	PROTEIN TRANSLOCATION,	MEMBRANE BOUND 3	PROTEINASE, MEMBRANE	TOWN TO THE TOWN	HYDROLASE SPASE I, LEADER PEPTIDASE I:	SERINE PROTEINASE.	SERINE-DEPENDANT	HYDROLASE, SIGNAL 2	PEPTIDE PROCESSING,	PROTEIN TRANSLOCATION,	MEMBRANE BOUND 3	PROTEIN ASE, IMEMBRAINE PROTEIN	G.	COMPLEX (ZINC	(ZINC FINGER/DNA), ZINC	FINGER, DNA-BINDING	FROIDIN
Compound	CHAIN: B; DNA (5'- CHAIN: C;	SIGNAL PEPTIDASE I;	CHAIN: A, B, C, D;							SIGNAL PEPTIDASE I;	CHAIN: A, B, C, D;						•	OF CALANT PROPERTY A CITY I	SIGNAL PEPTIDASE I; CHAIN: A. B. C. D:						,			QGSR ZINC FINGER	PEPTIDE; CHAIN: A; DUPLEX	OLIGONUCLEOTIDE	ם החשים ים והי החשותים
SEQ FOLD score																															
PMF score		0.78					,			0.47		,						5,5	0.43									0.41			
Verify score		-0.24								-0.25								0,00	09:0									0.15			
Psi Blast		2.6e-15								1.6e-16					-			25.75	1.6e-20									1.5e-27			
END AA		105								127								125	33									206			
START AA		38								38			•					96	82									113			
CHAIN		Ą								၁			_ -					6	<u>a</u>									А			
PDB ID		1612								1612								15.15	7191									laih			
SEQ ID NO:		316								316								316	316						•			317			

Pable 5

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PDB annotation		COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA	INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC) FINGER, PROTEIN A	INTERACTION, PROTEIN	DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC	FINGER/DNA) ZINC FINGER, PROTFIN-DNA	INTERACTION, PROTEIN	DESIGN, 2 CRYSTAL	SINCTIONE, COMPLEY	COMPLEX (ZINC	FINGER/DNA) ZINC FINGER, F	PROTEIN-DNA INTERACTION PROTEIN	DESIGN, 2 CRYSTAL	STRUCTURE, COMPLEX	COMPT EV (7TMC)	FINGER/DNA) ZINC FINGER.	PROTEIN-DNA	INTERACTION, PROTEIN	DESIGN, 2 CRYSTAL	STRUCTURE, COMPLEX	COMPLEX (ZINC	FINGER/DNA) ZINC FINGER, F
Compound		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;		T A A T TATE	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN: CHAIN: C F G:			DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER PROTEIN: CHAIN: CHAIN:	() () () () () () () () () ()			DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER	FROIEIN; CHAIN: C, F, G;		•	DNA: CHAIN A B D E.	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;				DNA: CHAIN: A. B. D. E.	CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;
SEQ FOLD score												-										•			
PMF score		0.87			00:1			1.00	-				1.00		•			90	7.00					1.00	
Verify score		0.08		9,0	84.0			0.43					0.16		-			0.36	05.0	-				0.36	
Psi Blast		1.5e-44		7, 7	3.46-49			6.8e-50		_			1.2e-50					1 20 50	1.26-30	-	•			1.2e-50	
A END		506		7.00	5 24			262					230					210	010					346	, ¹ -
START AA		125		621	ect.			181	,				209					727	/57					265	
CHAIN		ပ		ļ	ر			၁					ບ					ر	ر					ပ	
PDB UD		1mey			I mey			Imey					Imey					12001	, mey					Imev	
SEQ ID NO:		317		212	716			317					317					217	/16					317	

Pable 5

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PDB annotation	INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER DIA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER,	PROTEIN-DNA	DESIGN, 2 CRYSTAL	STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER FINGER	PROTEIN-DNA	INTERACTION, PROTEIN	DESIGN, 2 CRYSTAL	STRUCTURE, COMPLEX	COMPLEX (ZINC	FINGER/DNA) ZINC FINGER,	PROTEIN-DNA	INTERACTION, PROTEIN	DESIGN, 2 CRYSTAL	STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC	FINGERDNA) ZINC FINGER,	FROI EIN-LINA	DESIGN OFFICE A	CTDITCHIBE COMPLEX	(ZINC FINGER/DNA)	COMPLEX (TRANSCRIPTION:	REGULATION/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA), KNA	TRANSCRIPTION F	INITIATION, ZINC FINGER
Compound		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;			DNA; CHAIN: A, B, D, E;	PROTEIN; CHAIN: C, F, G,				DNA; CHAIN: A. B. D. E.	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;					DNA; CHAIN: A, B, D, E;	CONSENSOS ZINC FINGER	FROI BLIN; CHALIN: C. F. G;				TFIIIA; CHAIN: A, D; 5S	RIBOSOMAL RNA GENE;	CHAIN: B, C, E, F;			
SEQ FOLD score						93.81																							
PMF score		1.00									1.00							0.95						0.99	·				
Verify score		0.50									0.45			•				0.01						-0.14					
Psi Blast		1e-50				1e-50					1.4e-47							1.7e-12					-	1.4e-37					
A END	·	374				375					400							178						299					}
START AA		293				293					321							151						154					
CHAIN		၁				၁					S							5						Ą					
20 E		lmey				Imey					Imey							lmey						11f6					
SEQ ID NO:		317				317					317				_			317				,		317					

		T		_			_														11	Ha.	. 1	_	-	<u> </u>	i i	-IL-II			H	18. 1			
PDB annotation	PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA)	REGULATION/DNA), RNA	POLYMERASE III, 2	TRANSCRIPTION	INITIATION, ZINC FINGER	CONDITION OF AMECUNICAL	REGULATION/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA), RNA	POLYMERASE III, 2	TRANSCRIPTION	INITIATION, ZINC FINGER	PROTEIN	COMPLEX (TRANSCRIPTION	KEGULATION/DNA) YING-	YANG 1; TRANSCRIPTION	INITIATION, INITIATOR	ELEMENT, YY1, ZINC 2	FINGER PROTEIN, DNA-	PROTEIN RECOGNITION, 3	COMPLEX (TRANSCRIPTION	REGULATION/DNA)	COMPLEX (TRANSCRIPTION	KEGULATION/DINA) IING-	I ANG I; I KANSCKIF I JOIN	TO THE TAX THE TOTAL OF	ELEMENT, I II, SINC 2	PROTRIN PROCESSION 3	COMPLEX (TRANSCRIPTION	REGIT ATION/DNA)	COMPLEX (TRANSCRIPTION)	REGULATION/DNA) YING-	YANG 1; TRANSCRIPTION	**** *** *** *** ** ** ** ** ** *** **
Compound		TFIIIA; CHAIN; A, D; 5S RIBOSOMAL RNA GENE; CHAIN; B, C, F, B.	CILDAN: D, C, L, I,				TOTAL CITABLE A 15. 60	I FILIA; CHALIN: A, D; 3S RIBOSOMAL RNA GENE:	CHAIN: B, C, E, F;						YY1; CHAIN: C; ADENO-	ASSOCIATED VIKUS PS	INITIATOR ELEMENT	DNA; CHAIN: A, B;						YY1; CHAIN: C; ADENO-	ASSOCIATED VIKUS F3	INITIALOR ELEMENT	DIA, CIPIN. A, D,					YY1; CHAIN: C; ADENO-	ASSOCIATED VIRUS P5	INITIATOR ELEMENT	DIVIS, CALIMIN (3) 29
SEQ FOLD score		99.41																																	
PMTF					,		700	0.80							1.00									0.1		•						1.00			
Verify Score							200	60.0							-0.01									0.32								0.23			
Psi Blast		3.4e-38			-		2 10 20	3.46-38							5.1e-34									9.9e-50				-				3.4e-35	-		
END AA		370					392	70C							234				•					797								262			
START AA		209					238	007							127									158								191			
CHAIN		Ą					4	¢							ပ						_			ပ								C			
PDB ID		1tf6					- 1	9							Inbd									lubd								Iubd			
SEQ ID NO:		317					317	/10			,				317									317		_						317			

Table

PDB annotation	ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA- PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING- YANG 1; TRANSCRIPTION INTIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA- PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATIONIONA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATOR ELMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3
Compound		YY1; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	YY1; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YY1; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;
SEQ FOLD score					
PMF score		1.00	1.00	1.00	1.00
Verify score		0.03	0.18	0.10	0.11
Psi Blast		2e-50	1.7e-34	1.7e-35	9.9e-51
AA BA		290	318	346	374
START AA		179	217	245	263
CHAIN		ر ن	O	<u></u>	υ U
PDB ID		1ubd	1ubd	1ubd	lubd
SEQ ID NO:	·	317	317	317	317

Table 5

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PDB annotation	COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-	YANG I; TRANSCRIPTION	INITIATION, INITIATOR	FINGER PROTEIN, DNA-	PROTEIN RECOGNITION, 3	COMPLEX (TRANSCRIPTION	COMPLEX (TRANSCRIPTION	REGULATION/DNA) YING-	YANG 1; TRANSCRIPTION	INITIATION, INITIATOR	ELEMENI, YYI, ZINC 2	FINGER PROTEIN, DNA-	COMPLEX (TRANSCRIPTION	REGULATION/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA) YING	INTERVATION INTERVATION		ENGRED PROTEIN DIA-	PROTRIN RECOGNITION 3	COMPLEX (TRANSCRIPTION	REGULATION/DNA)		- ·	i dam.	*	4ms/	CINCINI AIN WHITE	PROTEIN/DNA) FIVE-	FINGER GLI, GLI, ZINC FINGER COMPIEX MA-	
Compound		YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS P5	INITIATOR ELEMENT	DNA; CHAIN: A, B;			÷	VV1. CHAIN: C. ADENO.	ASSOCIATED VIRUS PS	INITIATOR ELEMENT	DNA; CHAIN: A, B;				-	YY1; CHAIN: C; ADENO-	ASSOCIATED VIRUS P5	DNA: CHARLA B.	DINA; CHAIN: A, B;	•				COMPLEX(TRANSCRIPTIO	N REGULATION/DNA)	TRAMTRACK PROTEIN	(TWO ZINC-FINGER	PEPTIDE) COMPLEXED	WILL LUKE 3 DINA LUKE 4	GLII: CHAIN: A: DNA:	CHAIN: C, D,	
SEQ FOLD score		91.94			,																											
PMF score								000	}							0.86								0.12					5	7.00		
Verify score				•				-0.12								-0.25								-0.12					8	9.9		
Psi Blast		9.9e-51						3 30-47								16-31								9.9e-20				-	36 70 7	0.06-33		
END AA		375						400	3					•		400	-							206					170	707		
START AA		267						208	}							301								150					193	55		
CHAIN		ວ						ن	·)							၁								Y						۲		
PDB U	<u>.</u>	Iubd						111bd								lubd			-					2drp			-		::50	1187		
SEQ ID NO:		317						317								317								317					212	317.		

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						_	_																Pa: .				n +	TE 25				PP		n	
PDB annotation	BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING	PROTEIN/DNA) FIVE-	FINGER GLI; GLI, ZINC	FINGER, COMPLEX (DNA-	COMPLEX ONA BINDING	COMPLES DIVA-BINDING	PROTEIN/DIA) FIVE-	FINGER GLI; GLI, ZINC	FINGER, COMPLEX (DNA-	BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING	PROTEIN/DNA) FIVE-	FINGER GLL; GLL, ZINC	FINGER, COMPLEX (DNA-BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING	PROTEIN/DNA) FIVE-	FINGER GLI; GLI, ZINC	FINGER, COMPLEX (DNA-	BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING	PROTEIN/DNA) FIVE	FINGER GLI; GLI, ZINC	FINGER, COMPLEX (DNA-	BUNDANG FROIEMNIDINA)							BLOOD COAGULATION	EGF, HYDROLASE, SERINE	PROTEASE	SURFACE PROTEIN MEROZOITE SURFACE
Compound		ZINC FINGER PROTEIN	GLII; CHAIN: A; DNA;	CIPALIN: C, D;		ZINC HINCHE PROTEIN	OIL CHANGEN FROIDEN	GLII; CHAIN: A; DINA;	CHAIN: C, D;			ZINC FINGER PROTEIN	GLII; CHAIN: A; DNA;	Chair; C, J;		ZINC FINGER PROTEIN	GLII; CHAIN: A; DNA;	CHAIN: C, D;			ZINC FINGER PROTEIN	GLII; CHAIN: A; DNA;	CHAIN: C, D;				COAGULATION FACTOR EGF-LIKE MODULE OF	BLOOD COAGULATION	FACTOR X (N-TERMINAL,	1APO 3 APO FORM) (NMR,	13 STRUCTURES) IAPO 4	FACTOR VII; CHAIN:	,		MEROZOITE SURFACE PROTEIN 1; CHAIN: A;
SEQ FOLD score				-		81.75	67:10			,		,		•																					
PMF score		0.47			-						100	0.87				0.86					-0.12					30	3					B. 1			-0.13
Verify score		0.03										0.23				-0.16					0.21						7/.0				9,0	200			0.15
Psi Blast		1.7e-34				1.7e-34	5				3	0.86-34				1.7e-30					1.7e-30					3.	0.06-12				1, 2,	1.36-11			2.3e-09
END AA		317				348	}				220	3/3				366					205					5	<u>ئ</u>					70			78
START AA	·	189				209	}				2,75	242				273					65					5	3				5	3			22
CHAIN		A				A	:				ŀ	∢				¥					∀												•		V
PDB ID		2gli		_	_	2gli	ò				1	ng,				2gli					2gli					1	odar				641	Flor			Icej
SEQ ID NO:		317				317					217	710				317			•		317					220	026				320	920			320

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PDB annotation	ANTIGEN 1, MAJOR BLOOD- STAGE EGF-LIKE DOMAIN, EXTRACELLULAR, MODULAR PROTEIN, SURFACE 2 ANTIGEN, MALARIA VACCINE COMPONENT, SURFACE PROTEIN	BLOOD COAGULATION, SERINE PROTEASE, COMPLEX, CO-FACTOR, 2 RECEPTOR ENZYME, INHIBITOR, GLA, EGF, 3 COMPLEX (SERINE PROTEASE/COPACTOR/LIG AND)		BLOOD CLOTTING FACTOR VII, BLOOD COAGULATION, EGF-LIKE DOMAIN, BLOOD	BLOOD CLOTTING COMPLEX(SERINE PROTEASB/COFACTOR/LIG AND), BLOOD COAGULATION, 2 SERINE PROTEASB, COMPLEX, CO-PACTOR, RECEPTOR ENZYME, 3 INHIBITOR, GLA, BGF, COMPLEX (SERINB 4 PROTEASB/COFACTOR/LIG AND), BLOOD CLOTTING	PLASMINOGEN ACTIVATION
Compound		BLOOD COAGULATION FACTOR VIIA; CHAIN: L, H; SOLUBLE TISSUE FACTOR; CHAIN: T, U; D- PHE-PHE-ARG- CHLOROMETHYLKETONE (DFFRCMK) WITH CHAIN: C;	GROWTH FACTOR EPIDERMAL GROWTH FACTOR (EGF) (NMR, 16 STRUCTURES) 1EGF 3	BLOOD COAGILATION FACTOR VII, CHAIN: A;	BLOOD COAGULATION FACTOR VILA; CHAIN: L; BLOOD COAGULATION FACTOR VILA; CHAIN: H; SOLUBLE TISSUE FACTOR; CHAIN: T; 5L15; CHAIN: I;	T-PLASMINOGEN ACTIVATOR PI-G; 1TPG 7 CHAIN: NULL; 1TPG 8
SEQ FOLD score						
PMF		96:0	0.40	0.77	0.64	0.99
Verify score		0.30	0.18	<i>19</i> .0	0.42	0.45
Psi Blast		3.3e-10	2.6e-09	1.3e-11	3.3e-10	3e-11
END AA		78	89	69		29
START AA		23	26	23	23	22
CHAIN		1		¥	L	
PDB ID		1dan	legf	1 <i>f</i> 7e	lfak	14pg
SEQ ID NO:		320	320	320	320	320

Table 5

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PDB annotation	GLYCOPROTEIN GLYCOPROTEIN, HYDROLASE, SERINE PROTEASE, PLASMA, BLOOD 2 COAGULATION FACTOR	GROWTH FACTOR TGF- ALPHA, H-TGF-ALPHA; EGF- LIKB DOMAIN STRUCTURE, GROWTH FACTOR	SUGAR BINDING PROTEIN C-TYPE LECTIN, CRD, SP-D, COLECTIN, ALPHA- HELICAL COILED- 2 COIL, LUNG SURFACTANT, SUGAR BINDING PROTEIN	NK CELL NK CELL, RECEPTOR, C-TYPE LECTIN, C-TYPE LECTIN-LIKE, NKD	NK CELL NK CELL, RECEPTOR, C-TYPE LECTIN, C-TYPE LECTIN-LIKE, NKD	COLLAGEN BINDING PROTEIN IX-BP; IX-BP; COAGULATION FACTOR IX- BINDING, HETERODIMER, VENOM, HABU 2 SNAKE, C- TYPE LECTIN SUPERFAMILY, COLLAGEN BINDING PROTEIN	COLLAGEN BINDING PROTEIN IX-BP; IX-BP; COAGULATION FACTOR IX- BINDING, HETERODIMER, VENOM, HABU 2 SNAKE, C- TYPE LECTIN SUPERFAMILY, COLLAGEN BINDING PROTEIN
Compound	COAGULATION FACTOR X; CHAIN: NULL;	TRANSFORMING GROWTH PACTOR ALPHA; CHAIN: NULL;	LUNG SURFACTANT PROTEIN D; CHAIN: A, B, C;	CD94; CHAIN: NUIL;	CD94; CHAIN: NULL;	COAGULATION FACTOR IX-BINDING PROTEIN A; CHAIN: A; COAGULATION FACTOR IX-BINDING PROTEIN B; CHAIN: B;	COAGULATION FACTOR IX-BINDING PROTEIN A; CHAIN: A; COAGULATION FACTOR IX-BINDING PROTEIN B; CHAIN: B;
SEQ FOLD score					67.41	61.40	
PMF score	86.0	-0.07	0.93	1.00			0.99
Verify	0.92	0.50	69:0	0.32			0.42
Psi Blast	3.3e-10	2.3e-09	2.3e-22	1.6e-23	1.6e-23	9.9e-20	9.9e-20
END	,	99	372	376	376	374	372
START	23	56	242	249	249	249	250
CHAIN			A			∢ .	V
PDB US	Iwhe	lyuf	1508	1666	1b6e	1bj3	1bj3
SEQ ID NO:	320	320	321	321	321	321	321

Table 5

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PDB annotation	MEMBRANE PROTEIN C. TYPE LECTIN-LIKE DOMAINS	MEMBRANE PROTEIN C- TYPE LECTIN-LIKE DOMAINS	HEMATOPOIETIC CELL RECEPTOR ACTIVATION INDUCER MOLECULE (AIM), BA 1, HEMATOPOIETIC CELL RECEPTOR, LEUCOCYTE, C-TYPE LECTIN-LIKE, 2 NKD, KLR	SUGAR BINDING PROTEIN C-TYPE LECTIN, MANNOSE RECEPTOR	COAGULATION FACTOR BINDING IXX-BP COAGULATION FACTOR BINDING, Č-TYPE LECTIN, GLA-DOMAIN 2 BINDING, C- TYPE CRD MOTIF, LOOP EXCHANGED DIMER	COAGULATION FACTOR BINDING IXX-BP COAGULATION FACTOR BINDING, C-TYPE LECTIN, GLA-DOMAIN 2 BINDING, C- TYPE CRD MOTIF, LOOP EXCHANGED DIMER	COAGULATION FACTOR BINDING IXX-BP COAGULATION FACTOR BINDING, C-TYPE LECTIN, GLA-DOMAIN 2 BINDING, C- TYPE CRD MOTIF, LOOP EXCHANGED DIMER
Compound	FLAVOCETIN-A: ALPHA SUBUNIT; CHAIN: A; FLAVOCETIN-A: BETA SUBUNIT; CHAIN: B	FLAVOCETIN-A: ALPHA SUBUNIT; CHAIN: A; FLAVOCETIN-A: BETA SUBUNIT; CHAIN: B	EARLY ACTIVATION ANTIGEN CD69; CHAIN: A;	MACROPHAGE MANNOSE RECEPTOR; CHAIN: A, B;	COAGULATION FACTORS IXX-BINDING PROTBIN; CHAIN: A, B, C, D, E, F;	COAGULATION FACTORS IX/X-BINDING PROTEIN; CHAIN; A, B, C, D, B, F;	COAGULATION FACTORS IX/X-BINDING PROTEIN; CHAIN: A, B, C, D, E, F;
SEQ FOLD score		·			59.78		70.03
PMF score	0.81	0.98	1.00	0.75		1.00	
Verify score	0.61	0.36	0.91	12'0		0.38	
Psi Blast	6.6e-21	6.6e-22	1.6e-21	6.6e-22	2e-20	2e-20	6.6e-23
END AA	374	375	376	375	374	372	376
START	247	250	249	250	249	250	249
CHAIN	V	æ	∢ .	ø	V	Ą	<u>α</u>
PDB	lc3a	lc3a	1e87	legg	lixx	lixx	lixx
SEQ ID	321	321	321	321	321	321	321

Table 5

	·			_		٠,										- 312	PF-1	F	1.	F	E. 1L 1	· ·	- 11		L III	re fire	ar-tons
PDB annotation	COAGULATION FACTOR BINDING IX/X-BP	BINDING, C-TYPE LECTIN,	GLA-DOMAIN 2 BINDING, C-	EXCHANGED DIMER	LECTIN SUB-MBP-C; IRDL 9 C-TYPE LECTIN, CALCIUM-	BINDING PROTEIN IRDL 20	Process Property	KRINGLE 4, C-TYPE LECTIN,	2 CARBOHYDRATE RECOGNITION DOMAIN	LECTIN TETRANECTIN,	PLASMINOGEN BINDING,	KRINGLE 4, C-TYPE LECTIN,	2 CARBOHYDRATE RECOGNITION DOMAIN			et.	,	and?	INSECT IMMUNITY INSECT	HOMOPHILIC ADHESION	INSECT IMMUNITY INSECT	IMMUNITY, LPS-BINDING, HOMOPHII IC A DHESTON	GROWTH BACTOR/GROWTH	FACTOR RECEPTOR FGF,	FGFR, IMMUNOGLOBULIN-	LIKE, SIGNAL	DIMERIZATION, GROWTH F
Compound	COAGULATION FACTORS IX/X-BINDING PROTEIN;	CIPALIN: A, B, C, D, E, F;			MANNOSE-BINDING PROTEIN-C; IRDL 6	CHAIN: 1, 2; IRDL 7	IEIKANECIIN; CHAIN:	,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,		TETRANECTIN; CHAIN:	NULL;		·	LECTIN MANNOSE-	BINDING PROTEIN A	(LECTIN DOMAIN)	COMPLEX WITH 2MSB 3	CALCLIUM AIND GLYCOPEPTIDE 21MSB 4	HEMOLIN; CHAIN: A, B;		HEMOLIN; CHAIN: A, B;		HIRRORI AST GROWTH	FACTOR 2; CHAIN: A, B;	FIBROBLAST GROWTH	FACTOR RECEPTOR 1;	CERAIN: C, D;
SEQ FOLD score					53.97	20.03	07:70							55.49							·						
PMF score	0.63								<u></u>	1.00		•							0.04	•	82.0		0.70	2			
Verify score	0.62									0.86									-0.03		0.13		-0.03				
Psi Blast	6.6e-23				1.3e-18	6 60 22	77-20:0			6.6e-22				6.6e-19					9.9e-20		1.3e-23		1.36-18				
END	375				375	375	9/6			376				375					240		215		211	!)		•	
START AA	250				762	245	3			249				797					75		36		23	1	_		
CHAIN ID	В				-									¥					∀		A		U				
PDB ID	lixx				ligi	1rn3	2			1tn3				2msb					lbih 		1bih		lcvs				
SEQ ID NO:	321				321	321	125			321				321					325		325		325				

Table 5

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PDB annotation	PACTOR/GROWTH FACTOR RECEPTOR	GROWTH PACTOR/GROWTH FACTOR RECEPTOR FGF, FGFR, IMMUNOGLOBULIN-	LIKE, SIGNAL	TRANSDUCTION, 2	FACTOR/GROWTH PACTOR	RECEPTOR	GROWTH FACTOR/GROWTH	FACTOR RECEPTOR FGF,	LIKE, SIGNAL	TRANSDUCTION, 2	DIMERIZATION, GROWTH	FACTOR/GROWTH FACTOR RECEPTOR	GROWTH FACTOR/GROWTH	FACTOR RECEPTOR FGF1;	FGFR1; IMMUNOGLOBULIN	(IG) LIKE DOMAINS	BELONGING TO THE I-SET 2	SUBGROUP WITHIN IG-LIKE	IMMINE SYSTEM	MEMBRANE PROTEIN CD32;	FC RECEPTOR,	IMMUNOGLOULIN,	LEUKOCYTE, CD32	CONTRACTILE PROTEIN	BETA BARRET	Marca a portar	CONNECTIN, NEXTINS;	CELL ADHESION,	GLYCOPROTEIN,	I KANSMEMBKANE, REPEAT, BRAIN, 2
Compound		FIBROBLAST GROWTH FACTOR 2; CHAIN: A, B; FIBROBLAST GROWTH	FACTOR RECEPTOR 1;	CHAIN: C, D;			FIBROBLAST GROWTH	FACTOR 2; CHAIN: A, B;	FACTOR RECEPTOR 1;	CHAIN: C, D;			FIBROBLAST GROWTH	FACTOR 1; CHAIN: A, B;	FIBROBLAST GROWTH	FACTOR RECEPTOR 1;	CHAIN: C, D;		PC RECEPTOR	FC(GAMMA)RIIA; CHAIN:	A;	,		TELOKIN; CHAIN: A		TPETAL CHANG MITT.	inny, Chama, Molla,			
SEQ FOLD score															·															
PMF		0.18					0.39						0.18						0.24					0.64		76.0	5			
Verify score		-0.17					-0.15			-			-0.19				-		9.11					0.17		220	¥			
Psi Blast		3.3e-12	-				1.7e-22						3.36-20						3.3e-20					9.9e-12		2 20 12	C1-5C-7			
END		120					215						215						208					121		121	7			
START AA		13					23						23						22					23		0	<u> </u>			
CHAIN ID	·	Q					A						၁						A					¥			• .			
PDB ID		lcvs					lcvs						levt						1fcg	9				1fhg		1201	1			
SEQ ID NO:		325					325						325						325					325		325	3			·

Table 5

			· ·					
PDB annotation	IMMUNOGLOBULIN FOLD, ALTERNATIVE SPLICING, SIGNAL, 3 MUSCLE PROTEIN		GLYCOPROTEIN CD4; IMMUNOGLOBULIN FOLD, TRANSMEMBRANE, GLYCOPROTEIN, T-CELL, 2 MHC LIPOPROTEIN, POLYMORPHISM	MUSCLE PROTEIN IMMUNOGLOBULIN SUPERFAMILY, I SET, MUSCLE PROTEIN	IMMUNE SYSTEM P58 NATURAL KILLER CELL RECEPTOR; KIR, NATURAL KILLER RECEPTOR, INHIBITORY RECEPTOR, 2 IMMUNOGLOBULIN	IMMUNE SYSTEM P58 NATURAL KILLER CELL RECEPTOR; KIR, NATURAL KILLER RECEPTOR, INHIBITORY RECEPTOR, 2 IMMUNOGLOBULIN	IMMUNE SYSTEM CD32; RECEPTOR, FC, CD32, IMMUNE SYSTEM	CELL ADHESION PROTEIN F NCAM MODULE 2; CELL ADHESION, GLYCOPROTEIN, HEPARIN-
Compound		MUSCLE PROTEIN TITIN MODULE MS (CONNECTIN) 1TNM 3 (NMR, MINIMIZED AVERAGE STRUCTURE) 1TNM 4 1TNM 58	T-CELL SURFACE GLYCOPROTEIN CD4; CHAIN: A, B;	TWITCHIN 18TH IGSF MODULE; CHAIN: NULL;	MHC CLASS I NK CELL RECEPTOR PRECURSOR; CHAIN: A;	MHC CLASS I NK CELL RECEPTOR PRECURSOR; CHAIN: A;	FC GAMMA RIB; CHAIN: A;	NEURAL CELL ADHESION MOLECULE, LARGE ISOFORM; CHAIN: A;
SEQ FOLD score							•	
PMF score		0.17	0.59	0.13	0.24	0.36	0.21	0.12
Verify score		0.15	0.26	0.35	-0.13	0.07	-0.03	0.46
Psi Blast		3.36-11	9.9e-19	6.6e-14	1.3e-10	6.6e-22	3.3e-24	6.6e-11
END AA		121	223	121	120	209	209	120
START AA		36	33	24	14	42	24	25
CHAIN			Ą		Ā	A	Ą	Ą
PDB 13		1tmm	lwio	Iwit	2dli		2fcb	3ncm
SEQ ID NO:	·	325	325	325	325	325	325	325

Table :

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PDB annotation	BINDING, GPI-ANCHOR, 2 NBURAL ADHESION MOLECULE, INMUNOGLOBULIN FOLD, HOMOPHILIC 3 BINDING, CELL ADHESION PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DINA) COMPLEX (ZINC FINGER/DINA), ZINC FINGER, DINA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX
Compound		OGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B,	OGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B,	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B,	QOSR ZINC FINGER PEPTIDE; CHAIN: A;
SEQ FOLD score				70.01			
PMF		0.80	0.99	·	0.69	0.96	0.23
Verify score		-0.17	-0.12		0.00	-0.06	-0.16
Psi Blast		1.7e-30	6.6e-39	2c-39	2e-39	1.6e-38	1.6e-40
END		220		250	277	302	361
START AA	·	150	168	168	196	224	252
CHAIN		V	∢ .	∢ .	∢ .	₹	V
PDB ED		laih	lalh	laih	laih	lath	lalh
SEQ ID NO:		329	329	329	329	329	329

PDB annotation	(ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZÎNC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN		
Compound	DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCL EOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	DNA-BINDING PROTEIN HUMAN ENHANCER- BINDING PROTEIN MBP-1 MUTANT WITH CYS 11 1BBO 3 REPLACED BY ABU (C11ABU) (NMR, 60 STRUCTURES) 1BBO 4	DNA-BINDING PROTEIN HUMAN ENHANCER- BINDING PROTEIN MBP-1 MUTANT WITH CYS 11
SEQ FOLD score		·					
PMF score		96.0	1.00	0.92	0.76	0.03	0.37
Verify		0.25	0.02	-0.18	-0.15	-0.59	-0.40
Psi Blast		9.96-42	6.6e-44	3e-40	1.6e-38	6.6e-17	1.36-24
END AA		. 386	416	444	473	196	420
START AA		310	336	364	392	153	366
CHAIN	-	¥	¥	¥	∢		
20g ED ED		laih	laih	lalh	laih	1bbo	1bbo
SEQ ID NO:	·	329	329	329	329	329	329

Table

PDB annotation			COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER,DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX	COMPLEX (ZINC FINGER,DNA) ZINC FINGER,, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (TRANSCRIPTION
Compound	IBBO 3 REPLACED BY ABU (C11ABU) (NMR, 60 STRUCTURES) 1BBO 4	DNA-BINDING PROTEIN HUMAN ENHANCER- BINDING PROTEIN MBP-1 MUTANT WITH CYS 11 IBBO 3 REPLACED BY ABU (C11ABU) (NMR, 60 STRUCTURES) 1BBO 4	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, B; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	YY1; CHAIN: C; ADENO-
SEQ FOLD score							
PMF score		0.10	1.00	99.0	0.77	0.88	0.94
Verify score		-0.78	0.36	0.20	-0.37	0.39	0.04
Psi Blast		3.3e-23	6.6e-14	3.3e-13	3.3e-12	1.6e-12	3.3e-40
END		448	220	416	.200	528	248
START AA		394		389	473	205	150
CHAIN			ڻ ن	_ნ	ტ	ි. ජ	2
PDB ID		1950	Imey	Imey	Imey	Imey	Iubd
SEQ ID NO:	`	329	329	329	329	329	329

Table 5

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PDB annotation	REGUL, ATION/DNA) YING- YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA- PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGUL, ATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 ** COMPLEX (TRANSCRIPTION) REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING- YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA- PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION) REGULATION/DNA) YING- YANG 1; TRANSCRIPTION INITIATION, INITIATIOR
Compound	ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;
SEQ FOLD score				83.88	
PMF		1.00	1.00		0.40
Verify		0.09	-0.16		-0.17
Psi Blast		3.36-49	9.96-52	9.9e-52	6.6e-53
END		276	305	305	416
START AA		172	<u> </u>	<u>195</u>	277
CHAIN		ບ	U	U	ပ ပ
PDB ID		lubd	1ubd	lubd	lubd
SEQ ID NO:		329	329	329	329

Table 5

			·	Harry Versie and the Control of the	7	
PDB annotation	ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA- PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING- YANG I; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YYI, ZINC 2 FINGER PROTEIN, DNA- PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING- YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA- PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATOR BLEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 (COMPLEX (TRANSCRIPTION) REGULATION/DNA)		
Compound		YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	COMPLEX(TRANSCRIPTIO N REGULATION/DNA) TRAMTRACK PROTEIN (TWO ZINC-FINGER PEPTIDE) COMPLEXED WITH 2DRP 3 DNA 2DRP 4	COMPLEX (TRANSCRIPTIO
SEQ FOLD score					·	
PMF		0.65	0.89	1.00	0.35	0.49
Verify		-0.26	-0.50	-0.06	-0.17	0.35
Psi Blast	·	6.6e-51	9.96-52	3.36-53	9.9e-26	9.9e-28
AA.		472	501	228	220	302
START		361		418	164	248
CHAIN		ပ	O	ပ	A	Y
PDB UD		lubd	lubd	1ubd	2drp	2drp
SEQ ID		329	329	329	329	329

Table 5

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PDB annotation					COMPLEX (INHIBITOR/NUCLEASE) COMPLEX (INHIBITOR/NUCLEASE), COMPLEX (RI-ANG), HYDROLASE 2 MOLECULAR RECOGNITION, EPITOPE MAPPING, LEUCINE-RICH 3 REPEATS	COMPLEX (NUCLBAR PROTEIN/RNA) COMPLEX (NUCLEAR PROTEIN/RNA), (RNA, SNRNP, RIBONUCLEOPROTE)	COMPLEX (NUCLEAR PROTEINRNA), F (NUCLEAR PROTEIN/RNA), F RNA, SIRNP, RIBONUCLEOPROTEIN IN
Compound	N REGULATIONDNA) TRAMTRACK PROTEIN (TWO ZINC-FINGER PEPTIDE) COMPLEXED WITH 2DRP 3 DNA 2DRP 4	COMPLEX(TRANSCRIPTIO N REGULATIONDNA) TRAMTRACK PROTEIN (TWO ZINC-FINGER PEPTIDE) COMPLEXED WITH 2DRP 3 DNA 2DRP 4	COMPLEX(TRANSCRIPTIO N REGULATION/DNA) TRAMTRACK PROTEIN (TWO ZINC-FINGER PEPTIDE) COMPLEXED WITH 2DRP 3 DNA 2DRP 4	, and the second	RIBONUCLEASE INHIBITOR; CHAIN: A, D; ANGIOGENIN; CHAIN: B, E;	U2 RNA HAIRPIN IV; CHAIN: Q, R; U2 A; CHAIN: A, C; U2 B"; CHAIN: B, D;	UZ RNA HAIRPIN IV; CHAIN: Q, R; UZ A; CHAIN: A, C; UZ B'; CHAIN: B, D;
SEQ FOLD score							
PMF score		0.05	0.51		86.0	-0.01	96'0
Verify score		0.17	0.48		0.20	0.23	0.05
Psi Blast		1.6e-27	1.6e-29		6.6e-32	6.6e-16	1.6e-16
END AA		360	388		374	298	374
START		276	332		23	136	249
CHAIN	·	ď	4		∢	∢	¥
PDB U		2drp	2drp		1a4y	la9n	la9n
SEQ ID NO:		329	329		332	332	332

			·				
PDB annotation	COMPLEX (NUCLEAR PROTEIN/RNA) COMPLEX (NUCLEAR PROTEIN/RNA), RNA, SNRNP,RIBONUCLEOPROTE	IN COMPLEX (NUCLEAR PROTEIN/RNA) COMPLEX (NUCLEAR PROTEIN/RNA), RNA, SNRNP,RIBONUCLEOPROTE IN	COMPLEX (NUCLEAR PROTEINRNA) COMPLEX (NUCLEAR PROTEINRNA), RNA, SNRNP, RIBONUCLEOPROTE IN	COMPLEX (NUCLEAR PROTEIN/RNA) COMPLEX (NUCLEAR PROTEIN/RNA), RNA, SNRNP,RIBONUCLEOPROTE	COMPLEX (NUCLEAR PROTEIN/RNA) COMPLEX (NUCLEAR PROTEIN/RNA), RNA, SNRNP,RIBONUCLEOPROTE IN	GROWTH FACTOR/GROWTH FACTOR RECEPTOR FGF, FGFR, IMMUNOGLOBULIN- LIKE, SIGNAL TRANSDUCTION, 2 DIMERIZATION, GROWTH FACTOR/GROWTH FACTOR	GROWTH PACTOR/GROWTH
Compound	U2 RNA HAIRPIN IV; CHAIN: Q, R; U2 A; CHAIN: A, C; U2 B"; CHAIN: B, D;	U2 RNA HAIRPIN IV; CHAIN: Q, R; U2 A'; CHAIN: A, C; U2 B"; CHAIN: B, D;	UZ RNA HARPIN IV; CHAIN: Q. R; U2 A; CHAIN: A, C; U2 B"; CHAIN: B, D;	UZ RNA HAIRPIN IV; CHAIN: Q, R; U2 A; CHAIN: A, C; U2 B"; CHAIN: B, D;	UZ RNA HAIRPIN IV; CHAIN: Q. R; UZ A; CHAIN: A, C; UZ B"; CHAIN: B, D;	FIBROBLAST GROWTH FACTOR 2; CHAIN: A, B; FIBROBLAST GROWTH FACTOR RECEPTOR 1; CHAIN: C, D;	FIBROBLAST GROWTH
SEQ FOLD score							
PMF score	0.16	0.51	0.90	-0.02	0.78	0.24	0.18
Verify score	-0.20	0.51	0.09	0.15	-0.00	-0.02	0.04
Psi Blast	9.9e-22	6.6e-21	6.6e-24	3.3e-17	1.6e-16	6.6e-19	9.9e-18
END	183	207	242	309	374	519	519
START AA	09	2	98	111	249	404	404
CHAIN	V	A	∢	ပ	υ _.	<u>်</u>	Д
PDB ID	1a9n	1a9n	la9n	1a9n	la9n	lcvs	Icvs
SEQ ID NO:	332	332	332	332	332	332	332

Table 5

			· 	ter the steer the second			
PDB annotation	PACTOR RECEPTOR FGF, FGFR, IMMUNOGLOBULIN- LIKE, SIGNAL TRANSDUCTION, 2 DIMERIZATION, GROWTH FACTOR/GROWTH FACTOR	GROWTH FACTOR/GROWTH FACTOR RECEPTOR FGF1; FGFR1; IMMUNOGLOBULIN (IG) LIKE DOMAINS BELONGING TO THE I-SET 2 SUBGROUP WITHIN IG-LIKE DOMAINS, B-TRBFOIL FOLD	CONTRACTILE PROTEIN IMMUNOGLOBULIN FOLD, BETA BARREL	MUSCLE PROTEIN CONNECTIN, NEXTM5; CELL ADHESION, GLYCOPROTEIN, TRANSMEMBRANE, REPEAT, BRAIN, 2 IMMUNOGLOBULIN FOLD, ALTERNATIVE SPLICING, SIGNAL, 3 MUSCLE PROTEIN		MUSCLE PROTEIN IMMUNOGLOBULIN SUPERFAMILY, I SET, MUSCLE PROTEIN	NERVE GROWTH PACTOR/TRKA COMPLEX
Compound	FACTOR 2; CHAIN: A, B; FIBROBLAST GROWTH FACTOR RECEPTOR 1; CHAIN: C, D;	FIBROBLAST GROWTH FACTOR 1; CHAIN: A, B; FIBROBLAST GROWTH FACTOR RECEPTOR 1; CHAIN: C, D;	TELOKIN; CHAIN: A	TITIN; CHAIN: NULL;	MUSCLE PROTEIN TITIN MODULE M5 (CONNECTIN) 1TNM 3. (NMR, MINIMIZED AVERAGE STRUCTURE) 1TNM 4 1TNM 58	TWITCHIN 18TH IGSF MODULE; CHAIN: NULL;	NERVE GROWTH PACTOR; CHAIN: V, W;
SEQ FOLD score			·				
PMF score		0.58	0.95	0.59	0.80	0.18	-0.12
Verify		0.13	0.55	0.29	0.53	0.53	0.05
Psi Blast		6.66-19	1.6e-19	3.3e-21	9.9e-21	2.6e-20	3.3e-18
END A		519	501	501	501	200	501
START AA		404	409	410	415	409	416
CHAIN		ပ	Ą				×
PDB ID		levt	1fhg	lnct	1 thús	1wit	1ww w
SEQ ID NO:		332	332	332	332	332	332

Table 5

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PDB annotation	BETA-NGF; COMPLEX, TRKA RECEPTOR, NERVE GROWTH FACTOR, CYSTEINE KNOT, 2 IMMUNOGLOBULIN LIKE DOMAIN, NERVE GROWTH	ACETYLATION RNASE INHIBITOR, RIBONUCLEASE/ANGIOGEN IN INHIBITOR ACETYLATION, LEUCINE- RICH REPEATS	CELL ADHESION PROTEIN NCAM MODULE 2; CELL ADHESION, GLYCOPROTEIN, HEPARIN- BINDING, GPI-ANCHOR, 2 NEURAL ADHESION MOLECULE, IMMUNOGLOBULIN FOLD, HOMOPHILIC 3 BINDING, CELL ADHESION PROTEIN	OXIDOREDUCTASE METHYLENETHF DEHYDROGENASB / METHENYLTHF THF, BIFUNCTIONAL, DEHYDROGENASE, CYCLOHYDROLASE, FOLATE, 2 OXIDOREDUCTASE HBADER	OXIDOREDUCTASE,HYDRO LASE FOLATE, DEHYDROGENASE, CYCLCOHYDROLASE, BIFUNCTIONAL, 2
Compound	TRKA RECEPTOR; CHAIN: X, Y;	RIBONUCLEASE INFIBITOR; CHAIN: NULL;	NEURAL CELL ADHESION MOLECULA, LARGE ISOFORM; CHAIN: A;	METHYLENETETRAHYDR OFOLATB DEHYDROGENASE / CHAIN: A, B;	FOLD BIFUNCTIONAL PROTEIN; CHAIN: A;
SEQ FOLD score				·	
PMF score		0.93	0.80	0.98	0.63
Verify score		0.31	0.33	0.33	0.22
Psi Blast	·	9.9c-44	2.36-20	0.00023	0.00066
END AA		361	200	511	511
START AA		09	410	403	412
CHAIN			V	V	Y
PDB ID		2bnh	Зпсш	1a4i	160a
SEQ ID NO:		332	332	333	333

Table 5

 -			г	- _T		12. 0	
PDB annotation	CHANNELING, OXIDOREDUCTASE,HYDRO LASE	OXIDOREDUCTASE GLUTAMIC DEHYDROGENASE; GLUTAMATE DEHYDROGENASE, ALLOSTERY, ABORTIVE COMPLEX	OXIDOREDUCTASB (CHOH(D)-NAD+(A)) R- LACTATE DEHYDROGENASE; 2DLD 7	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INTIATOR HEMBRY, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	CONTRACTILE PROTEIN TRIPLE-HELIX COILED COIL, CONTRACTILE PROTEIN	COMPLEX (GTP- BINDING/TRANSDUCER) BETA1, TRANSDUCIN BETA
Compound		GLUTAMATE DEHYDROGENASE; CHAIN: A, B, C, D, B, F;	D-LACTATE DEHYDROGENASE; 2DLD 5 CHAIN: A, B; 2DLD 6	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	HUMAN SKELETAL MUSCLE ALPHA-ACTININ 2; CHAIN: A;	GT-ALPHA/GI-ALPHA CHIMERA; CHAIN: A; GT- BETA; CHAIN: B; GT-
SEQ FOLD score							
PMF score		0.24	0.75	-0.01	0.01	0.00	0.27
Verify score		0.26	0.01	0.17	-0.14	-0.27	0.47
Psi Blast		0.002	1.3e-30	2e-16	1.36-16	9.9e-15	6.6e-14
END		517	594	180	180	1279	147
START AA			400	109	104	666	2
CHAIN		¥	V	<	U	A	æ
PDB CI		1ch6	2dld	laih	Iubd	Iquu	lgot
SEQ ID NO:	·	333	333	338	338	339	341

		•											_	-		<u> </u>		i	<u> </u>			· •	117	-	174	
PDB annotation	SUBUNIT; GAMMA1, TRANSDUCIN GAMMA SUBUNIT; COMPLEX (GTP- BINDING/TRANSDUCER), G PROTEIN, HETEROTRIMER 2 SIGNAL TRANSDUCTION		COMPLEX (KINASE/INHIBITOR) CDK6;	PI9INK4D; CYCLIN DEPENDENT KINASE,	CYCLIN DEPENDENT	PROTEIN, CDK, INK4, CELL	CYCLE, COMPLEX	(KINASE/INHIBITOR)	COMPLEX (INHIBITOR	PROTEIN/KINASE)	INHIBITIOR PROTEIN,	CYCLIN-DEPENDENT	KINASE, CELL CYCLE 2	CONTROL, ALPHA/BETA,	COMPLEX (INHIBITOR	PHOSPHOTE ANSFER ASP	PROTEIN KINASE ICKI 18		OXYGEN TRANSPORT	HEME PERPERATORY	PROTEIN, ERYTHROCYTE	OXYGEN TRANSPORT	OXYGEN TRANSPORT,	PROTEIN REVIEW !	OXYGEN TRANSPORT	OXYGEN TRANSPORT, HEME, RESPIRATORY
Compound	GAMMA; CHAIN: G;		CYCLIN-DEPENDENT KINASE 6; CHAIN: A, C;	CYCLIN-DEPENDENT KINASE INHIBITOR;	CHAIN: B, D;		•		CYCLIN-DEPENDENT	KINASE 6; CHAIN: A;	P19INK4D; CHAIN: B;					CACRIN KINASR I DRI TA.	ICKI 6 CHAIN: A, B; 1CKI 7		HEMOGLOBIN; CHAIN: A,	4		HEMOGLOBIN; CHAIN: A,	Д		HEMOGI OBIN: CHAIN: A	В
SEQ FOLD score																						135.78			102 38	
PMF			0.03			<u> </u>			0.24							0.17			1.00							,
Verify score			-0.23						-0.01							-0 33			0.65					٠.		
Psi Blast			66000.0						0.0023							0.0066			2e-47			2e-47			336-38	
A END			204						200							157			141		,	141			141	
START			83						83							3,4	3		1			-			-	•
CHAIN			Ą						A			-				V			A			Ą	٠.		и	1
PDB BDB			1bi8						Iblx							1.51			1a4f			la4f			1a4f	
SEQ ID NO:			342			,			342							342			344			344			344	

Table 5

				7						_						-proj.		P	P. CO.	- (1-1)			 - -	-p- :	<u></u>	
PDB annotation	PROTEIN, ERYTHROCYTE	OXYGEN TRANSPORT OXYGEN TRANSPORT, HEME, RESPIRATORY PROTEIN, ERYTHROCYTE	OXYGEN TRANSPORT OXYGEN TRANSPORT	OXYGEN TRANSPORT OXYGEN TRANSPORT													4	OXYGEN	HEME, OXYGEN DELIVERY	VEHICLE, BLOOD	SUBSTITUTE	OXYGEN TRANSPORT.	CHIMERA PROTEIN,	RESPIRATORY PROTEIN,	OVYCENT TO ANSBODT	OXYGEN TRANSPORT,
Compound		HEMOGLOBIN; CHAIN: A, B	HEMOGLOBIN; CHAIN: A, E, C, F;	HEMOGLOBIN; CHAIN: A, E, C, F;	OXYGEN TRANSPORT HEMOGLOBIN	THIONVILLE ALPHA	VAL 1 1BAB 3 REPLACED	BY GLU AND AN	BOUND TO THE 1BAB 4	AMINO TERMINUS 1BAB 5	OXYGEN TRANSPORT	THIONVILLE ALPHA	CHAIN MUTANT WITH	VAL 1 IBAB 3 REPLACED	BY GLU AND AN	ACELYLALED MEI BOUND TO THE 1BAB 4	AMINO TERMINUS 1BAB 5	DEOXYHEMOGLOBIN	A: DEOXYHEMOGLOBIN	(BETA CHAIN); CHAIN: B,	D;	CHIMERA HEMOGLOBIN	BETA-ALPHA; CHAIN: A,	B,C,D;	MODELLE STREETHER	CHIMERA HEMOGLOBIN
SEQ FOLD score		-	106.12		135.21										•						.0.0.	105.91				
PMF score		1.00		1.00							1.00					•		1.00							1 00	1.00
Verify score		0.85		0.79							0.78							0.89							07.0	0.72
Psi Blast		3.3e-38	1.3e-38	1.3e-38	9.9e-46						9.9e-46							3e-46			9	3.3e-42			2 22 40	3.36-42
END		140	140	140	141						141							141				141			121	141
START AA		2	1	2	1						7						,	1			•				,	7
CHAIN ID		Ø	Ħ	凹	Ą						∢							Ą				∢				∢
908 E0		la4f	la9w	la9w	Ibab						1bab							1c7c				1cn4			77.7	Ich4
SEQ ID NO:		344	344	344	344						344			,				344				\$			244	344

Table 5

PDB annotation	CHIMERA PROTEIN, RESPIRATORY PROTEIN, HEME			OXYGEN STORAGE/TRANSPORT HB D; HB D HEMOGLOBIN D (R- STATE) 1, HEMOGLOBIN,	COOPERATIITY, OXYGEN TRANSPORT	OXYGEN STORAGE/TRANSPORT HB D; HB D HEMOGLOBIN D (R-	STATE) I, HEMOGLOBIN, AVIAN, HIGH 2 COOPERATIITY, OXYGEN	STORAGETRANSPORT HB (STATE) I, HEMOGLOBIN, CAVIAN, HIGH 2 COOPERATIITY, OXYGEN	DE/		OXYGEN TRANSPORT X- II RAY STUDY, PORCINE HEMOGLOBIN, ARTIFICIAL [1]
Compound	BETA-ALPHA; CHAIN: A, B, C, D;	OXYGEN CARRIER HEMOGLOBIN (DEOXY) 1HBH 3	OXYGEN CARRIER HEMOGLOBIN (DEOXY) 1HBH 3	HEMOGLOBIN D; CHAIN: A, C; HEMOGLOBIN D; CHAIN: B, D;		HEMOGLOBIN D; CHAIN: A, C; HEMOGLOBIN D; CHAIN: B, D;		HEMOGLOBIN D; CHAIN: A, C; HEMOGLOBIN D;	CHALN: B, D;	OXYGEN TRANSPORT HEMOGLOBIN (DEOXY) 1HDA 3	OXYGEN TRANSPORT HEMOGLOBIN (DEOXY) IHDA 3	PORICINE HEMOGLOBIN (ALPHA SUBUNIT); CHAIN: A, C; PORICINE
SEQ FOLD score		120.38	,			148.19		96.34			136.61	
PMIF score	,		1.00	1.00	,		·			1.00		1.00
Verify score			0.61	0.93						0.49		0.78
Psi Blast		9.9e-45	9.9e-45	9.9e-45		9.9e-45		6.6e-38		3.3e-47	3.3e-47	3.3e-47
END AA		141	141	141		141		138		141	141	141
START AA		1	2	1		I	•	1		_	1	1
CHAIN	·	Ą	¥	V		¥		æ		¥	A	A
PDB U		મુવ્યા	11154	1hbr		1hbr		1hbr		1hda	1hda	lqpw
SEQ ID NO:		344	344	344		344		344		344	344	344

					<u> </u>												!	(lands	n	n		[] Serve	u 14.	ft een	<u>۔</u> م	. 19			er this	(Long)
PDB annotation	HUMAN BLOOD, 2 OXYGEN TRANSPORT	OXYGEN TRANSPORT X- RAY STUDY, PORCINE	HEMOGLOBIN, ARTIFICIAL	TRANSPORT	OXYCHN TP ANSPORT	OXYGEN TRANSPORT	HEME, RESPIRATORY	PROTEIN, ERYTHROCYTE	OXYGEN TRANSPORT	OXIGEN IRANSPORT,	PROTEIN, ERYTHROCYTE	OXYGEN TRANSPORT	OXYGEN TRANSPORT,	PROTEIN ERVTHROCYTE	OXYGEN TRANSPORT	HEME PROTEIN, MODEL	COMPOUNDS, OXYGEN	STORAGE, LIGAND 2	BINDING GEOMETRY,	CONFORMATIONAL	SUBSTATES, OXYGEN 3	OXYGEN TRANSPORT	OXYGEN IKANSPORT	tum	# * -	· Lina	de well	the Unit	dia Char	thata.
Compound	HEMOGLOBIN (BETA SUBUNIT); CHAIN: B, D	PORICINE HEMOGLOBIN (ALPHA SUBUNIT);	CHAIN: A, C; PORICINE	SUBUNIT); CHAIN: B, D	HEMOGI OBIN: CHAIN: A	B	:		HEMOGLOBIN; CHAIN: A,	20		HEMOGLOBIN; CHAIN: A.	m		MYOG! OBIN: CHAIN:	NULL:		_				HEMOGLOBIN; CHAIN: A,	E, C, F;	OXYGEN TRANSPORT	THIONVILLE ALPHA	CHAIN MUTANT WITH	VAL 1 1BAB 3 REPLACED	BY GLU AND AN	ACETYLATED MET	AMINO TERMINUS 1BAB 5
SEQ FOLD score		133.84							106.59		-	87.72										93.82								
PMF					0.08	97.0									0.49	:			•					1.00						
Verify			• .		0.03	3					•			_	0.49	}		_						0.38					•	
Psi Blast	-	3.3e-47			3 30-47	2			3.3e-42			2e-32			1.66-38							3e-33		26-40	•					_
END AA		141			170	:			179			179			179		,					178		179						
START AA		-			-	4			6			37			_							37		2						
CHAIN		¥			A	•			¥			В										B		∢						
PDB TO		Iqpw			19df				1a4f			1a4f			1a6m							la9w		1bab						
SEQ ID NO:		344			378	}			345			345			345							345		345						

Table 5

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PDB annotation			OXYGEN STORAGE/TRANSPORT HEME, OXYGEN DELIVERY VEHICLE, BLOOD SUBSTITUTE	OXYGEN TRANSPORT OXYGEN TRANSPORT, CHIMERA PROTEIN, RESPIRATORY PROTEIN, HEME	OXYGEN TRANSPORT OXYGEN TRANSPORT, CHIMERA PROTEIN, RESPIRATORY PROTEIN, HEMB	months these times (fear
Compound	OXYGEN TRANSPORT HEMOGLOBIN THIONVILLE ALPHA CHAIN MUTANT WITH VAL 1 IBAB 3 REPLACED BY GLU AND AN ACETYLATED MET BOUND TO THE IBAB 4 AMINO TERMINUS IBAB 5	OXYGEN TRANSPORT HEMOGLOBIN THIONVILLE ALPHA CHAIN MUTANT WITH VAL 1 IBAB 3 REPLACED BY GLU AND AN ACETYLATED MET BOUND TO THE IBAB 4 AMINO TERMINUS IBAB 5	DEOXYHEMOGLOBIN (ALPHA CHAIN); CHAIN: A; DEOXYHEMOGLOBIN (BETA CHAIN); CHAIN: B, D;	MODULE-SUBSTITUTED CHIMERA HEMOGLOBIN BETA-ALPHA; CHAIN: A, B, C, D;	MODULE-SUBSTITUTED CHIMERA HEMOGLOBIN BETA-ALPHA; CHAIN: A, B, C, D;	OXYGEN TRANSPORT MYOGLOBIN COMPLEXED WITH CYANIDE 1EMY 3 IEMY 107 HEME PROTEIN, GLOBIN FOLD 1EMY 5
SEQ FOLD score	113.33	86.57			94.07	
PMF score			0.25	0.99		0.95
Verify score			0.23	0.13		0.44
Psi Blast	2e-40	3.3e-35	6.6e-41	1.6e-36	1.6e-36	6.6e-38
END AA	179	179	179	6 <i>i</i> 1	179	179
START AA	37	43	proof.	7	37	2
CHAIN	V	m	¥	₹	V	
PDB U	1bab	1bab	1c7c	1ch4	Ich4	lemy
SEQ ID NO:	345	345	345	345	345	345

							19 2 5- 1	mente methor	t is there there	سور در پوست	ii ti saab	med ment
PDB annotation	·				OXYGEN — STORAGE/TRÂNSPORT HB D; HB D HEMOGLOBIN D (R-	SIAIE) I, HEMOGLOBIN, AVIAN, HIGH 2 COOPERATITY, OXYGEN TRANSPORT	OXYGEN STORAGE/TRANSPORT HB D; HB D HEMOGLOBIN D (R- STATE) 1, HEMOGLOBIN,	AVIAN, HIGH 2 COOPERATIITY, OXYGEN	OXYGEN STORAGE/TRANSPORT HB D; HB D HEMOGLOBIN D (R-	AVIAN, HIGH 2 COOPERATIITY, OXYGEN		grad grad times to the control of th
Compound	OXYGEN TRANSPORT HEMOGLOBIN (DEOXY, HUMAN FETAL F=/II\$=) IFDHG 1 IFDHH 2	OXYGEN CARRIER HEMOGLOBIN (DEOXY) 1HBH 3	OXYGEN CARRIER HEMOGLOBIN (DEOXY) 1HBH 3	OXYGEN CARRIER HEMOGLOBIN (DEOXY) 1HBH 3	HEMOGLOBIN D; CHAIN: A, C; HEMOGLOBIN D; CHAIN: B, D;		HEMOGLOBIN D; CHAIN: A, C; HEMOGLOBIN D; CHAIN: B, D;		HEMOGLOBIN D; CHAIN: A, C; HEMOGLOBIN D; CHAIN: B, D;		OXYGEN TRANSPORT HEMOGLOBIN (DEOXY) 1HDA 3	OXYGEN TRANSPORT HEMOGLOBIN (DEOXY)
SEQ FOLD score	100.14		101.11	76.75			117.48		81.42			105.99
. PMF score		0.36			1.00				·		1.00	
Verify score		0.16			0.75						0.49	
Psi Blast	1.3e-33	3.3e-39	3.3e-39	6.6e-36	3e-39	,	3e-39		3.3e-32		1.3e-41	1.3e-41
END AA	179	179	179	179	179		179		176		179	179
START AA	39	2	39	38	1		37	·	37			40
CHAIN	_ව	Ą	¥	В	¥		4		m		¥	¥
PDB TD	1fdh	lhbb	1hbh	1hbh	lhbr		lhbr		lhbr		lhda	1hda
SEQ ID NO:	345	345	345	345	345		345		345		345	345

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PDB annotation			OXYGEN TRANSPORT X-	HEMOGLOBIN ARTIFICIAL	HUMAN BLOOD, 2 OXYGEN	TRANSPORT	OXYGEN TRANSPORT X.	HEMOGLOBIN, ARTIFICIAL	HUMAN BLOOD, 2 OXYGEN	TRANSPORT	OXYGEN TRANSPORT X-	HEMOGLOBIN, ARTIFICIAL	HUMAN BLOOD, 2 OXYGEN	TRANSPORT	OXYGEN TRANSPORT	HBTNCO; HBTNCO;	OXYGEN TRANSPORT, HEMOGLORIN		COMPLEX (TRANSCRIPTION	FACTOR/DNA) BHLH;	UASP2(17); TRANSCRIPTION	FACTOR, BASIC HELLX	LOOP HELIX, 2 COMPLEX	(TRANSCRIPTION	FACTOR/DNA)	COMPLEX (TRANSCRIPTION	14. STEPOL DEGIT ATORY	ELEMENT BINDING	ELIX-	LOOP-HELIX-LEUCINE	ZIPPER, SREBP,	ווערוויים בייים בייים הייים ווערים ווערים בייים
Compound	1HDA 3	OXYGEN TRANSPORT HEMOGLOBIN (DEOXY) 1HDA 3	PORICINE HEMOGLOBIN	CHAIN: A. C. PORICINE	HEMOGLOBIN (BETA	SUBUNIT); CHAIN: B, D	PORICINE HEMOGLOBIN	CHAIN: A. C. PORICINE	HEMOGLOBIN (BETA	SUBUNIT); CHAIN: B, D	PORICINE HEMOGLOBIN (ALPHA SUBINITY):	CHAIN: A, C; PORICINE	HEMOGLOBIN (BETA	SUBUNIT); CHAIN: B, D	HEMOGLOBIN; CHAIN: A;	HEMOGLOBIN; CHAIN: B;			PHOSPHATE SYSTEM	POSITIVE REGULATORY	PROTEIN CHAIN: A, B;	DNA; CHAIN: C, D;				STEROL REGULATORY	PROTEIN 14. CHAIN: A B	C. D. DNA: CHAIN: E. F. G.	H;	•		Lane
SEQ FOLD score		82.30					110.57				88.40				80.38																•	
PMF score			1.00																0.55							90.0						
Verify score			0.46																-0.25							-0.46						
Psi Blast		3.3e-36	1.6e-41	***********			1.6e-41				2e-35				3e-36				0.00033							1.3e-07						
END AA		179	179	·			179				179				179				136							149						
START AA		6	1				9			į	37				38				83							83						
CHAIN		Д	Ą				∢			١	29				m				Ą							<		٠.				
PDB ED		1hda	Iqpw				Iqpw				Idpw				ltln				1a0a							lam9	•					1
SEQ ID NO:		345	345				345			2,50	345				345				348							348						

Table 5

	1.		· · · · · · · · · · · · · · · · · · ·	,						
PDB annotation	COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) SREBP-1A; STEROL REGULATORY ELEMENT BINDING PROTEIN, 2 BASIC-HELIX-LOOP-HELIX-LEUCINE ZIPPER, SREBP,	TRANSCRIPTION 3 FACTOR, COMPLEX (TRANSCRIPTION REGULATION/DNA)			METAL BINDING PROTEIN RING FINGER PROTEIN MAT1; RING FINGER (C3HC4)	T Tank Tank	TRANSFERASE HRS; HRS, VHS, FYVE, ZINC FINGER, SUPERHELIX	METAL BINDING PROTEIN RING FINGER PROTEIN MAT1; RING FINGER (C3HC4)	PATHOGENESIS-RELATED
Compound		STEROL REGULATORY ELEMENT BINDING PROTEIN 1A; CHAIN: A, B, C, D; DNA; CHAIN: E, F, G, H;		VIRUS EQUINE HERPES VIRUS-1 (C3HC4, OR RING DOMAIN) ICHC 3 (NMR, 1 STRUCTURE) 1CHC 4	VIRUS EQUINE HERPES VIRUS-1 (C3HC4, OR RING DOMAIN) ICHC 3 (NMR, 1 STRUCTURE) 1CHC 4	CDK-ACTIVATING KINASE ASSEMBLY FACTOR MAT1; CHAIN: A;	VIRUS EQUINE HERPES VIRUS-1 (C3HC4, OR RING DOMAIN) 1CHC 3 (NMR, 1 STRUCTURE) 1CHC 4	HEPATOCYTE GROWTH FACTOR-REGULATED TYROSINE CHAIN: A;	CDK-ACTIVATING KINASE ASSEMBLY FACTOR MAT1; CHAIN: A;	PATHOGENESIS-RELATED
SEQ FOLD score						·				84.39
PMF score		0.29		0.00	0.28	0.41	0.00	0.41	0.41	
Verify score		-0.55		-0.78	0.03	-0.76	-0.78	-0.44	-0.76	
Psi Blast		3.3e-07		0.0023	0.00099	2.3e-05	0.0023	0.0097	2.3e-05	3.3e-41
END		153		223	08	224	188	187	189	212
		8.		193	29	161	158	153	156	53
CHAIN START ID AÁ		ф				Ą		A	A	
PDB ID		lam9		lchc	1chc	1925	1chc	ldvp	1g25	1cfe
SEQ ID NO:		348		352	352	352	353	353	353	354

Table 5

		_		r	PET			
PDB annotation	PROTEIN PATHOGENESIS- RELATED LEAF PROTEIN 6, ETHYLENE PATHOGENESIS- RELATED PROTEIN, PR-1 PROTEINS, 2 PLANT	PATHOGENESIS-RELATED PROTEIN PATHOGENESIS- RELATED LEAF PROTEIN 6, ETHYLENE PATHOGENESIS- RELATED PROTEIN, PR-1 PROTEINS, 2 PLANT DEFENSE	PATHOGENESIS-RELATED PROTEIN PATHOGENESIS- RELATED LEAF PROTEIN 6, ETHYLENE PATHOGENESIS- RELATED PROTEIN, PR-1 PROTEINS, 2 PLANT	ALLERGEN ANTIGEN 5; ANTIGEN 5, ALLERGEN, VESPID VENOM	ALLERGEN ANTIGEN 5; ANTIGEN 5, ALLERGEN, VESPID VENOM	TRANSFERASB APS KINASB; APS KINASE, ADENYLYLSULFATE KINASE, SULFATE, NUCLEOTIDE 2 KINASE, TRANSFERASE		
Compound	PROTEIN P14A; CHAIN: NULL; 	PATHOGENESIS-RELATED PROTEIN P14A; CHAIN: NULL:	PATHOGENESIS-RELATED PROTEIN P14A; CHAIN: NULL;	VES V 5; CHAIN: A;	VES V 5; CHAIN: A;	ADENOSINE- SPHOSPHOSULFATE KINASE; CHAIN: A, B;	ZINC FINGER /DNA\$ BINDING DOMAIN ZINC FINGER (NMR\$) 3ZNF 3	HYDROLASE(O-
SEQ FOLD score								
PMF		66.0	0.95	1.00	1.00	0.25	0.13	0.21
Verify score		0:30	0.40	0.23	0.31	0.03	0.26	0.02
Psi Blast		5.1e-35	3.36-41	8.5e-39	3.3e-42	3.3e-05	0.00051	1.6e-15
END AA	-	212	212	212	213	28	172	154
START AA		53	95	24	47	17	143	81
CHAIN			·	¥	₹ .	Ą		
PDB UI		Icfe	lcfe	Iqnx	Iqnx	146j	3znf	1191
SEQ ID NO:	·	354	354	354	354	357	359	360

Table 5

		· · · · · · · · · · · · · · · · · · ·	Time start respectively and the start start of start at the start
PDB annotation			Cfan
Compound	GLYCOSYL) LYSOZYME (B.C.3.2.1.17) MUTANT WITH CYS 54 REPLACED BY THR, 119L 3 CYS 97 REPLACED BY ALA, ALA 134 REPLACED BY SER (C54T,C97A, 119L 4 A134S) 119L 5	HYDROLASE(O- GLYCOSYL) LYSOZYME (B.C.3.2.1.17) MUTANT WITH CYS 54 REPLACED BY THR, 119L 3 CYS 97 REPLACED BY ALA, ALA 134 REPLACED BY SER (C54T,C97A, 119L 4 A134S) 119L 5	HYDROLASE (O-GLYCOSYL) LYSOZYME (E.C.3.2.1.17) MUTANT WITH THR 34 REPLACED BY ALA, 174L 3 LYS 35 REPLACED BY ALA, SER 36 REPLACED BY ALA, PRO 37 174L 4 REPLACED BY ALA, SER 38 REPLACED BY 174L 5 ALA, SER 38 REPLACED BY 174L 5 ALA, SER 4R REPLACED BY ALA, SER 8 REPLACED BY 174L 5 ALA, CLU 45 REPLACED BY 174L 7 REPLACED BY 174L 7 ALA, CYS 54 REPLACED BY 174L 7 THR, CYS 54 REPLACED BY 174L 7 REPLACED BY 174L 7 ALA ALA CYS 97 REPLACED BY ALA ALA ALA (T34A,K35A,S36A,P37A,S38 D,N40A, 174L 8
SEQ FOLD score			
PMF score		0.11	0.03
Verify score		-0.12	-0.09
Psi Blast		5.16-11	1.40-11
END AA	·	203	203
START		83	≅
CHAIN			¥
PDB U		1191	1741
SEQ ID NO:		360	360

| Table

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PDB annotation							,																							-				, 1			
Compound	S44A,E45A,D47A,K48A,C54 T,C97A) 174L 9	HYDROLASE (O-	GLYCOSYL) LYSOZYME	WITH LEU 32 REPLACED	BY THR, 176L 3 THR 34	REPLACED BY LYS, LYS	35 REPLACED BY VAL,	SER 36 176L 4 REPLACED	BY ASP, PRO 37	REPLACED BY GLY, SER	38 REPLACED BY 176L 5	ASN, LEU 39 REPLACED	BY SER, CYS 54	REPLACED BY THR, AND	176L 6 CYS 97 REPLACED	BY ALA	(L32T,T34K,K35V,S36D,P37	G,S38N, 176L 7	L39S,C54T,C97A) 176L 8	HYDROLASE (O-	GLYCOSYL) LYSOZYME	(E.C.3.2.1.17) MUTANT	WITH LEU 32 REPLACED	BY THR, 176L 3 THR 34	REPLACED BY LYS, LYS	35 REPLACED BY VAL,	SER 36 176L 4 REPLACED	BY ASP, PRO 37	REPLACED BY GLY, SER	38 REPLACED BY 176L 5	ASN, LEU 39 REPLACED	BY SER, CYS 54	REPLACED BY THR, AND	176L 6 CYS 97 REPLACED	BY ALA	(L321,134K,K33 V,S36L,F37 G,S38N, 176L 7	
SEQ FOLD score	•		•																																		
PMF score		0.40																		0.03																	
Verify score		-0.04																		0.10																	
Psi Blast		1.6e-15																		1.7e-12																	
END AA		154																		203																	
START AA		80																	·	81																	
CHAIN		A	•			•												-		∢		_											,				
PDB UD		1761																		1761																	
SEQ ID NO:		360																		360																	

Table 5

					-	7		_				_			_	_	_	7	.	P	-	- 4	-		Į.	5 (e i	T.		E		
PDB annotation															HYDROLASE (O-	GLYCOSYL)	HYDROLASE (O-	GLYCOSYL)	HYDROLASE (O- GLYCOSYL)												LIGASE PHERS; PHERS;	ENZYME, TRNA	HOMODIMER	
Compound	L39S,C54T,C97A) 176L 8	HYDROLASE (O-	GLYCUSYL) LYSUZYME (E.C.3.2.1.17) MUTANT	WITH ILE 3 REPLACED BY	LEU, 189L 3 SER 38	KEPLACED BY ASP, ALA	41 REFLACED BI VAL. AI A 82 1891 4 PRO ASN	116 REPLACED BY ASP.	VAL 131 REPLACED BY	ALA, 189L 5 AND ASN 144	REPLACED BY ASP	SUBSTITUTIONS	(I3L,S38D,A41V, 189L 6	A82P,N116D,V131A,N144D) 1891.7	LYSOZYME; 191L 4	CHAIN: NULL; 191L 5	LYSOZYME; 191L 4	CHAIN: NULL; 191L S	LYSOZYME; 192L 4 CHAIN: NULL: 192L 5	HYDROLASE(Ó-	GLYCOSYL) LYSOZYME	(E.C.3.2.1.17) INSERTION	MUTANT WITH ALA ALA	ALA 205L 3 INSERTED	AFTER SER 44, CYS 54	REPLACED BY THR, CYS	97 205L 4 REPLACED BY	ALA (INS(S44-	AAA),C54T,C97A) 205L 5		PHENYLALANYL-TRNA	SYNTHETASE; CHAIN: A;	SYNTHETASE, CHAIN: B;	
SEQ FOLD score																			!															
PMF		0.35													0.51		0.21		0.03	0.43											00.0			
Verify		-0.01									,		•		0.11		0.13		-0.28	0.15											-0.38			
Psi Blast		1.3e-15													1.3e-16		1.7e-11		3.4e-11	le-11				•							5.1e-52	•		
A EN		154													154		203		203	203									ļ		416		:	
START		80													81		81		81	81							,				2			
CHAIN																															В			
PDB ID		1891						- -							1161		1911		1921	2051											1b7y			
SEQ ID NO:		360													360		09E		360	360											361			

Table 5

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PDB annotation	TRANSCRIPTION REGULATION PROTO- ONCOGENE, NUCLEAR BODIES (PODS), LEUKEMIA, 2 TRANSCRIPTION REGULATION	TRANSCRIPTION REGULATION PROTO- ONCOGENE, NUCLEAR BODIES (POD\$), LEUKEMIA, 2 TRANSCRIPTION REGULATION			METAL BINDING PROTEIN RING FINGER PROTEIN MAT1; RING FINGER (C3HC4)	DNA-BINDING PROTEIN V(D)J RECOMBINATION ACTIVATING PROTEIN 1; RAGI, V(D)J RECOMBINATION, ANTIBODY, MAD, RING FINGER, 2 ZINC BINUCLEAR CLUSTER, ZINC FINGER, DNA-BINDING PROTEIN		45cc
Compound	TRANSCRIPTION FACTOR PML; CHAIN: NULL;	TRANSCRIPTION FACTOR PML; CHAIN: NULL;	VIRUS EQUINE HERPES VIRUS-1 (C3HC4, OR RING DOMAIN) ICHC 3 (NMR, 1 STRUCTURE) 1CHC 4	VIRUS EQUINE HERPES VIRUS-1 (C3HC4, OR RING DOMAIN) ICHC 3 (NMR, 1 STRUCTURE) ICHC 4	CDK-ACTIVATING KINASE ASSEMBLY FACTOR MAT1; CHAIN: A;	RAGI; CHAIN: NULL;	VRUS EQUINE HERPES VRUS-1 (C3HC4, OR RING DOMAIN) ICHC 3 (NMR, 1 STRUCTURE) ICHC 4	VIRUS EQUINE HERPES
SEQ FOLD score								
PMF score	0:00	0.04	0.62	1.00	0.07	0.70	0.84	0.84
Verify	-0.89	-0.59	0.19	-0.01	-0.27	80.0-	-0.07	-0.07
Psi Blast	2.36-12	1.7e-05	6.6e-16	1.7e-14	6.6e-14	1.4e-07	6.8e-05	9.96-06
A END	338	332	342	333	338	332	155	155
	282	283	288	289	288	284	109	109
CHAIN					⋖			
PDB U	1bor	1bor	Ichc	1chc	1825	1md	lchc	1chc
SEQ ID NO:	364	364	364	364	364	364	365	365

Table 5

		77	g			FL 1905	PE	/Uli	
ton		METAL BINDING PROTEIN RING FINGER PROTEIN MAT1; RING FINGER (C3HC4)	DNA-BINDING PROTEIN V(D)J RECOMBINATION ACTIVATING PROTEIN 1; RAG1, V(D)J RECOMBINATION, ANTIBODY, MAD, RING FINGER, 2 ZINC BINUCLEAR CLUSTER, ZINC FINGER, DNA-BINDING PROTEIN	TPR- L HSP70,		COMPLEX OF TWO ELONGATION FACTORS EF- TU; EF-TS; ELONGATION FACTOR, NUCLEOTIDE EXCHANGE, GTP-BINDING, 2 COMPLEX OF TWO ELONGATION FACTORS	RNA BINDING PROTEIN G- PROTEIN, BETA-BARREL	TRANSLATIONAL GIPASE EF-G RIBOSOMAL TRANSLOCASE, TRANSLATIONAL GIPASE	RNA BINDING PROTEIN I EFTU; TRANSPORT AND II PROTECTION PROTEIN, RNAI
PDB annotation		DING PROTEINGE	NG PRC MBINA G PRO1 I ATION, MAD, I INC BII INC FIN	E HOP, EPTIDE HELICA IC70, 21		OF TWO ON PAC SLONG, SCLEO', GTP-B	VG PRO ETA-B/	ONAL OMAL ASE, ONAL	NG PRO NSPORT N PRO
PDB		METAL BINDING PROT RING FINGER PROTEIN MAT1; RING FINGER (C3HC4)	DNA-BÍNDING PROTEIN V(D)I RECOMBINATION ACTIVATING PROTEIN I; RAGI, V(D)I RECOMBINATION, ANTIBODY, MAD, RING FINGER, 2 ZINC BINUCLE CLUSTER, ZINC FINGER, DNA-BINDING PROTEIN	CHAPERONE HOP, TPR- DOMAIN, PEPTIDE- COMPLEX, HELICAL REPEAT, HSC70, 2 HSP70, PROTEIN BINDING		COMPLEX OF TWO ELONGATION PACTORS TU; EF-TS; ELONGATION FACTOR, NUCLEOTIDE EXCHANGE, GTP-BINDIN 2 COMPLEX OF TWO ELONGATION FACTORS	RNA BINDING PROTEIN (PROTEIN, BETA-BARREL	TRANSLATIONAL EF-G RIBOSOMAL TRANSLOCASB, TRANSLATIONAL	RNA BINDING PROTEIN EFTU; TRANSPORT AND PROTECTION PROTEIN,
		METAL RING FI MAT1; F	ANT ANT ANT PINC CLU	CHA CON CON REPU			PRO PRO	RA H RA	RNA BET
	REING JAMR, 1 4	AIN: A;	3	.D. 'Q.		ELONGATION FACTOR TU; CHAIN: A, B, E, F; ELONGATION FACTOR TS; CHAIN: C, D, G, H;	OR TU B, C, D	OR G.	OR;
Compound	VIRUS-1 (C3HC4, OR RING DOMAIN) 1CHC 3 (NMR, 1 STRUCTURE) 1CHC 4	CDK-ACTIVATING KINASE ASSEMBLY FACTOR MATI; CHAIN: A;	RAGI; CHAIN: NULL;	TPR1-DOMAIN OF HOP; CHAIN: A, B; HSC70- PEPTIDE; CHAIN: C, D;		ELONGATION FACTOR TU; CHAIN: A, B, E, F; ELONGATION FACTOR CHAIN: C, D, G, H;	ELONGATION FACTOR TU (EF-TU); CHAIN: A, B, C, D	ELONGATION FACTOR G; CHAIN: NULL;	ELONGATION FACTOR; CHAIN: A, B;
S	IS-1 (C3 AIN) 1C ICTURE	CDK-ACTIVATING KINASE ASSEMBLY FACTOR MAT1; CH	I; CHAI	-DOMA N: A, B IDE; CF		ELONGATION FAC TU; CHAIN: A, B, E ELONGATION FAC CHAIN: C, D, G, H;	GATIO U; CH	ELONGATION CHAIN: NULL;	ELONGATIO CHAIN: A, B;
	VIRU DOM STRU	RINA PACI	RAG	TPR1 CHAI PEPT		CHAN CHAN	ELO EL	CHA	CHA
SEQ FOLD score									
SEO							· 		
PMF score		0:30	0.09	9.68		-0.18	-0.15	-0.07	-0.15
· Verify score		-0.14	-0.76	-0.28		0.07	0.40	0.18	0.12
Psi Blast		3.3e-06	0.001	5.1e-07		1.7e-38	1.4e-33	1,4e-12	3.46-41
END		152	186	39			219	155	219
START		117	102	: E		8	20	=	20
CHAIN		A		¥		4	A		A
PDB	,	1g25	Irmd	lelw		laip	1d2e	ldar	lefc
SEQ ID NO:									
SE		365	365	367	. '	369	369	369	369

Table 5

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PDB annotation	BINDING PROTEIN	-	HYDROLASE ERA, GTPASE, RNA-BINDING, RAS-LIKE, HYDROLASE			GTPASE, MOLECULAR	SWITCH, TRNA, RIBOSOME,	CHAPERONE, DISULFIDE ISOMER ASE	TRANSLATION EF-G; BENT	DOMAIN III, MUTATION	HIS573ALA	TRANSLATION TRANSLATIONAL GIPASE	PROTEIN BINDING EF-G; EF-	TRANSLOCASE, RIBOSOME,	ELONGATION, 2	TRANSLATION, PROTEIN	SYNT FACTOR, GTPASE,	UCLEOTIDE	BINDING, PROTEIN	DINCHAR	COMPLEX (ZINC FINGER/DINA) COMPLEX (ZINC FINGER/DINA), ZINC
Compound			GTP-BINDING PROTEIN ERA; CHAIN: A, B;	TRANSPORT AND PROTECTION PROTEIN ELONGATION FACTOR TU (DOMAIN I) - *GUANOSINE DIPHOSPHATE 1ETU 4	COMPLEX 1ETU 5	ELONGATION FACTOR TO (FF-TU): CHAIN: A:			ELONGATION FACTOR G;	Circles, 7s,		TRANSLATION INITIATION PACTOR IF2/EIF5B; CHAIN: A;	ELONGATION FACTOR G	CHAIN: A; ELUNGALIUN PACTOR G DOMAIN 3:	CHAIN: B;			•			QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX
SEQ FOLD score																					
PMF			-0.12	-0.14		-0.12			-0.14			-0.09	-0.14	-							0.23
Verify			0.12	0.21		0.09			0.17			0.23	0.19								-0.28
Psi Blast			3.4e-35	1.7e-37		1.7e-43			1.7e-12			6.8e-18	1.7e-12								9.9e-42
END			225	208		220			155			213	155		-						262
START			25	16		16			19			21	19								153
CHAIN			Ą			∢_			A			A	Ą						,		A
PDB			lega	letu		lexm			1fnm			1g7s	2efg								lalh
SEQ ID			369	369		369			369			369	369								370

Table 5

PDB annotation	FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN
Compound	OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDB; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;
SEQ FOLD score		81.16					
PMF score			0.17	1.00	1.00	1.00	1.00
Verify			-0.13	0.40	0.42	0.30	0.47
Psi Blast		3.36-37	3.36-37	9.9e-36	6.6e-37	3.3e-36	1.6e-36
A EN		319	346	430	457	486	513
START AA		237	237	349	377	405	433
CHAIN		V	V	¥	V	¥	V
PDB		laih	lalh	laih	lalh	lalh	laih
SEQ ID NO:		370	370	370	370	370	370

Table 5

PDB annotation	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, F PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL
Compound	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN; A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN; C, F, G;
SEQ FOLD score					·	·
PMF score	0.13	1.0	1.00	1.00	1.00	1.00
Verify score	0.20	0.49	0.55	0.44	0.38	0.08
Psi Blast	1.7e-25	1.76-44	6.8e-46	le-46	1.2e-47	1.2e-48
END	177	205	233	261	289	317
START	97	124	152	180	208	236
CHAIN	∢	U	ပ	ပ	ပ	ပ
EDB EDB	laih	Imey	Imey	Imey	Imey	1mey
SEQ ID NO:	370	370	370	370	370	370

Table 5

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PDB annotation	STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN, DNA	INTERACTION, PROTEIN	DESIGN, 2 CRYSTAL	STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER.	PROTEIN-DNA	INTERACTION, PROTEIN	STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC FINGER, FINGER,	PROTEIN-DNA	INTERACTION, PROTEIN	DESIGN, 2 CRYSTAL	STRUCTURE, COMPLEX	COMPLEX (ZINC	FINGER/DNA) ZINC FINGER	PROTEIN-DNA	INTERACTION, PROTEIN	DESIGN, 2 CRYSTAL	STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC	FINGER/DNA) ZINC FINGER	PROTEIN-DNA	INTERACTION, PROTEIN	DESIGN, 2 CRYSTAL	STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER FI	TINOTAL PARTY CONTRACTOR OF THE PARTY OF THE
Compound		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN: CHAIN: C B G:	incitation canalists of the			DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;				DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;				DNA: CHAIN: A B D E:	CONSENSUS ZINC PINGER	PROTEIN; CHAIN: C, F, G;				DNA; CHAIN: A. B. D. E.	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G,				DNA; CHAIN: A, B, D, B;	CONTRACTOR THAT A MACHINA
SEQ FOLD score		107.60									·																		
PMF score						1.00		•	,		1,00					8	7.00					1.00						1.00	
Verify score				·		0.34					9:00				•	032	70.0					0.47						99.0	
Psi Blast		1.5e-49				1.5e-49					1.46-50					170.50	7.76-30					3.4e-50						3.4e-50	
END		318				345					373					401	1					429						457	
START AA		236			•	264					262					320	020					348						376	
CHAIN		၁	-			၁				•	C.					ر	ر ر					U						U	
PDB ED		lmey				Imey					1mey					1mev	TIMES					lmev	,					lmey	
SEQ ID NO:		370				370					370					370	2					370						370	1

PDB annotation	PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CR YSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX	COMPLEX (ZINC FINGER, FINGER, DNA ZINC FINGER, FROTEIN-DNA INTERACTION, PROTEIN II DESIGN, 2 CR YSTAL STRUCTURE, COMPLEX II (ZINC FINGER, DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER F PROTEIN-DNA INTERACTION, PROTEIN IL DESIGN, 2 CRYSTAL
Compound	PROTEIN; CHAIN: C, P, G;	DNA; CHAIN: A, B, D, E, CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E, CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E, CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, B; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;
SEQ FOLD score	·				·	
PMF score		1.00	1.00	00.1	0.36	1.00
Verify score		0.40	0.53	0.46	-0.01	0.54
Psi Blast		5.1e-50	6.8e-50	1.4e-33	5.1e-42	3.4e-11
END AA		485	513	517	177	177
START		404	432	460	96	150
CHAIN		ပ	ပ	υ	ن د	O
PDB UD		Imey	Imey	lmey	lmey	Imey
SEQ ID NO:		370	370	370	370	370

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PDB annotation	(ZINC FINGER/DNA)	COMPLEX (ZINC	FINGER/DNA) ZINC FINGER, PROTFIN-DNA	INTER ACTION PROTEIN	DESIGN 2 CRYSTAL	STRICT TO THE STREET	(ZINC FINGER)DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA), RNA	POLYMERASE III, 2	TRANSCRIPTION	INITIATION, ZINC FINGER	PROTEIN	COMPLEX (TRANSCRIPTION	REGULATION/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA), RNA	POLYMERASE III, 2	TRANSCRIPTION	INTITATION, ZINC FINGER	PROTEIN	COMPLEX (TRANSCRIPTION	REGULATION/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA), RNA	POLYMERASE III, 2	TRANSCRIPTION	INITIATION, ZINC FINGER	PROTEIN	COMPLEX (TRANSCRIPTION,	REGULATION/DNA)	COMPLEX (IRANSCRIPTION	KEGULATION/DNA), KNA	TOLIMENABLIII, & III	IRANSCRIPTION INITIATION, ZINC FINGER FI
Compound		DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER PROTEIN: CHAIN:					TFIIIA; CHAIN: A, D; 5S	RIBOSOMAL RNA GENE:	CHAIN: B, C, B, F;						TFIIIA; CHAIN: A, D; 5S	RIBOSOMAL RNA GENE;	CHAIN: B, C, E, F;						TFIIIA; CHAIN: A, D; 5S	RIBOSOMAL RNA GENE;	CHAIN: B, C, E, F;						TFIIIA; CHAIN: A, D; 5S	KIBOSOMAL RNA GENE;	CHAIN: B, C, B, F;			
SEQ FOLD score								114.40			•																								-		·
PMF score		1.00											٠.			1.00								1.00								1.00					
Verify score		0.54														0.41								0.18		•						0.03					
Psi Blast		6.6e-15	····				,	5.1e-37								1.4e-35								5.1e-37								1.5e-38					
AA END		177						287			٠.					270								326								410					
START		150						122								125								181								265					
CHAIN		Ö						¥								∢								Ą								∢					-
PDB ED		lmey					, -	1tf6								1tte								1tf6								1066					
SEQ ID NO:		370						370								370								370								370					

PDB annotation	PROTEIN	COMPLEX (TRANSCRIPTION PROTICTION ATTOMINAL)	COMPLEX (TRANSCRIPTION	REGULATIONDNA), RNA	TRANSCRIPTION	INITIATION, ZINC FINGER	PROTEIN	COMPLEX (TRANSCRIPTION REGIT ATTOMONA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA), RNA	POLYMERASE III, 2	TRANSCRIPTION	INITIATION, ZINC FINGER	PROTEIN	COMPLEX (TRANSCRIPTION	REGULATION/DNA) YING-	YANG 1; TRANSCRIPTION	INITIATION, INITIATOR	ELEMENT, YY1, ZINC 2	PROTEIN RECOGNITION, 3 📆	RIPTION	REGULATION/DNA)	COMPLEX (TRANSCRIPTION,	REGULATION/DNA) YING-	YANG I; TRANSCRIPTION	INITIALION, INITIALOR	ELEMENT, YY1, ZINC2	PROTEIN PECOCNITION 3	COMPLEX (TRANSCRIPTION)	REGULATION/DNA)	COMPLEX (TRANSCRIPTION		INITIATION, INITIATOR II
Compound		THIIA; CHAIN: A, D; 5S	CHAIN: B, C, E, F;					THILL; CHAIN: A, D; 5S RIBOSOMAL RNA GENE:	CHAIN: B, C, E, F;						YY1; CHAIN: C; ADENO-	ASSOCIATED VIRUS PS	INITIATOR ELEMENT	DNA; CHAIN: A, B;					YY1; CHAIN: C; ADENO-	ASSOCIATED VIRUS PS	INITIATOR ELEMENT	LINA; CHAIN! A, B;					YY1; CHAIN: C; ADENO-	ASSOCIATED VIRUS PS	INITIATOR ELEMENT DNA; CHAIN: A, B;
SEQ FOLD score						•							-																				
PMF score		0.95						1.00	•						1.00								1.00								1.00		
Verify score		0.17						0.50					•		0.42								0.30								0.44		
Psi Blast		1.7e-37						3.4e-36							3.4e-31								6.6e-53								3.4e-32		
END		495						515							233								261							•	261		
START AA		349						377							127								127								160		
CHAIN		Ą						∢							ပ်	-				 			ပ							. ,	ပ		
PDB ID		1tf6						1#6							1ubd								Inbd								1 ubd		
SEQ ID NO:		370						370							370								370								370		

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PDB annotation	ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA- PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATIONDNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING- YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA- PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING- YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA- PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING- YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 (INCOMPLEX (TRANSCRIPTION) REGULATION/DNA)	COMPLEX (TRANSCRIPTION, REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INTIATION HELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-FINGER PROTEIN, DNA-FINGER PROTEIN, DNA-FINGER PROTEIN, DNA-FINGER PROTEIN, DNA-FINGER PROTEIN, DNA-FINGER PROTEIN, DNA-FINGER PROTEIN, DNA-FINGER PROTEIN, DNA-FINGER PROTEIN, DNA-FINGER PROTEIN, DNA-FINGER PROTEIN, DNA-FINGER PROTEIN RECOGNITION, 3 FINGER PROTEIN RECOGNITION RECOGNITION RECOGNITION RECOGNITION RECOGNITION RECOGNITION
Compound	ELF FIN PRC COJ		YY1; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 RECINITATOR ELEMENT YAI DNA; CHAIN: A, B; ELA FIN FIN FIN FIN FIN FIN FIN FIN FIN FIN	YY1; CHAIN: C; ADENO-COI ASSOCIATED VIRUS P5 REGINITIATOR ELEMENT YA DNA; CHAIN: A, B; RITE INT FIN FIN FIN FIN FIN FIN FIN FIN FIN FIN	YYI; CHAIN: C; ADENO- CO ASSOCIATED VIRUS P5 REC INITIATOR ELEMENT YA DNA; CHAIN: A, B; ELI FIN
SEQ FOLD score		YY1 ASS INIT DNA	YYI. ASSK INII DNA	YY1 ASS(INIT DNA	YYI ASSA INTI DNA
PMF score		1.00	1.00	1.00	1.00
Verify	·	0.15	0.22	0.23	0.50
Psi Blast		3.36-46	5.1e-33	3.4e-34	1.4e-35
END		318	317	345	373
START		213	216	42	266
CHAIN	·	ပ	v	O	υ
PDB U		lubd	1ubd	lubd	Iubd
SEQ ID		370	370	370	370

Table 5

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PDB annotation	COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING- VANG 1: TRANSCRIPTION	INITIATION, INITIATOR	ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-	PROTEIN RECOGNITION, 3	COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-	YANG 1; TRANSCRIPTION	INITIATION, INITIATION	FINGER PROTEIN, DNA-	PROTEIN RECOGNITION, 3	COMPLEX (TRANSCRIPTION	ACAMETER (TER ANSCHIEGTON)	REGIL ATTONONA) YING-	YANG 1: TRANSCRIPTION	INITIATION, INITIATOR	ELEMENT, YY1, ZINC 2	FINGER PROTEIN, DNA-	PROTEIN RECOGNITION, 3	REGULATIONONA)	COMPLEX (TRANSCRIPTION)	REGULATION/DNA) YING-	INTERIOR INTERIOR	BLEMENT, YY1, ZINC 2	FINGER PROTEIN, DNA-	PROTEIN RECOGNITION, 3	COMPLEX (TRANSCRIPTION)	COMPLEX (TRANSCRIPTION)
Compound		YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOD BI BMENT	DNA; CHAIN: A, B;				YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5	INITIATOR ELEMENT	DNA; CHAIN: A, B;				CITIES OF STREET	I I; CHALIN: C; AUDINO-	INITIATOR ELEMENT	DNA; CHAIN: A, B;	•		. `		YY1; CHAIN: C; ADBNO-	ASSOCIATED VIRUS P5	INITIALOR ELEMENT	DINA, CHAIN. A, D,				YY1; CHAIN: C; ADENO-
SEQ FOLD score		96.52																										
PMF score			. · ·				1.00							3.							1.00		-					1.00
Verify score							0.05						3	0.20						,	90.0							0.46
Psi Blast		1.3e-48					1.3e-48						}	2.16-30							1.3e-47							5.1e-35
END AA		402					402							104							458		٠					457
START		294					297							3						٠	346	,						356
CHAIN		ပ					ပ							۔ ر							၁							ပ
PDB U		lubd					lubd							pont							1ubd							lubd
SEQ ID		370					370							3/0							370		•					370

Table 5

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PDB annotation	REGULATION/DNA) YING- YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA- PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING- YANG I; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YYI, ZINC 2 FINGER PROTEIN, DNA- PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 TO COMPLEX (TRANSCRIPTION-PREGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION [] ELMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-ROTEIN RECOGNITION, 3; COMPLEX (TRANSCRIPTION) REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING- [I] YANG 1; TRANSCRIPTION [I] INITIATION, INITIATOR [I]
Compound	ASSOCIATED VIRUS P5 INTIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO-ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;
SEQ FOLD score					
PIMIF score		66:0	1.00	00'1	1.00
Verify score	· · · · · · · · · · · · · · · · · · ·	0.15	0.16	90.04	0.18
Psi Blast		6.66-46	5.1e-35	1.7e-34	1.7e-25
END		513	485	513	517
START AA		374	381	412	440
CHAIN		ပ	ပ	ပ	ပ
PDB U		Tubd	1ubd	lubd	lubd
SEQ ID		370	370	370 .	370

Table 5

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PDB annotation	ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA- PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE- FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE- FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEINDNA) FIVE- FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLJ; GLJ, ZINC FINGER, COMPLEX (DNA-FINGER, COMPLEX (DNA-FINGER) FINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTBIN/DNA) FIVB-FINGER GILI; GILI, ZINC FINGER, COMPLEX (DNA-FINDING PROTEIN/DNA)	TRANSPORT PROTEIN TC4; US GIPASE, NUCLEAR TRANSPORT, TRANSPORT	TRANSPORT PROTEIN TC4; TGTPASE, NUCLEAR TRANSPORT, TRANSPORT [1] PROTEIN
Compound		ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	GTP-BINDING PROTEIN RAN; CHAIN: A, B;	GTP-BINDING PROTEIN RAN; CHAIN; A, B;
SEQ FOLD score			99.97				55.66	
PMF score		1.00		1.00	66'0	0.64		1.00
Verify score		0.58		0.29	0.45	0.09		0.62
Psi Blast		1.4e-31	3.46-34	3.4e-34	1.5e-34	1.7e-29	1.7e-48	1.7e-48
END AA		260	319	3 4 4	512	207	186	181
START AA		124	180	216	384	88	19	20
CHAIN ID		٧	∀	¥	V V	V	A	æ
PDB UD		2gli	2gli	2gli	2gli	2gli	1byu	lbyu
SEQ ID NO:		370	370	370	370	370	371	371

Table 5

PROTIO-ONKOGENE SERMETHREONINE PROTEIN KINASE CHAIN: B; RAS-RELATED PROTEIN RAP-14; CHAIN: A; PROTO-ONKOGENE SERINE/THREONINE PROTEIN KINASE CHAIN: B; TRANSFORMING PROTEIN P21/H-RAS-1; CHAIN: A; PROTEIN KINASE CHAIN: A; PROTEIN KINASE CHAIN: A; PROTEIN KINASEORMING PROTEIN RHOA(0-181); CHAIN: A; PKN; CHAIN: B; RAB6 GTPASE; CHAIN: A; ELONGATION FACTOR G; CHAIN: NULL;		HADTEN-PROTEIN E COMPLEX, BIFFECTORS THAIN: SIGNALING PROTEIN GTP- BINDING PROTEIN BECOMPLEX, EFFECTORS THAIN: COMPLEX, EFFECTORS THAIN: ROTEIN, GTP HYDROLYSIS, KINETIC CRYSTALLOGRAPHY, 2 SIGNALING PROTEIN G N: A; HYDROLYSIS, KINETIC CRYSTALLOGRAPHY, 2 SIGNALING PROTEIN ROTEIN, GTP HYDROLYSIS, KINETIC CRYSTALLOGRAPHY, 2 SIGNALING PROTEIN SIGNALING PROTEIN SIGNALING PROTEIN SIGNALING PROTEIN TRAFFICKING TRAFFICKING TRAFFICKING TRAFFICKING TRAFFICKING TRAFFICKING TRAFFICKING TRAFFICKING TRAFFICKING TRAFFICKING TRAFFICKING TRAFFICKING TRANSLATIONAL GTPASE, RAB6, VESICULAR TRAFFICKING TRANSLATIONAL GTPASE TRANSLATIONAL GTPASE TRANSLATIONAL GTPASE TRANSLATIONAL GTPASE TRANSLATIONAL GTPASE TRANSLATIONAL GTPASE TRANSLATIONAL GTPASE TRANSLATIONAL GTPASE TRANSLATIONAL GTPASE TRANSLATIONAL GTPASE TRANSLATIONAL GTPASE
RAP-1A; CHAIN: A; RAP-1A; CHAIN: A; RAP-1A; CHAIN: A; PROTO-ONKOGENE SERINE/THREONINE PROTEIN KINASE CHAIN: B; TRANSFORMING PROTEIN P21/H-RAS-1; CHAIN: A; TRANSFORMING PROTEIN P21/H-RAS-1; CHAIN: A; FKIOA(0-181); CHAIN: A; PKN; CHAIN: B; RAB6 GTPASE; CHAIN: A; CHAIN: NULL; CHAIN: NULL;		
RAS-RELATED PROTEI RAP-1A; CHAIN: A; PROTO-ONKOGENE SERINE/THREONINE PROTEIN KINASE CHA B; TRANSFORMING PROT P21/H-RAS-1; CHAIN: A HIS-TAGGED TRANSFORMING PROT RANSFORMING PROT RHOA(0-181); CHAIN: A PKN; CHAIN: B; RAB6 GTPASE; CHAIN: CHAIN: NULL;	RAS-RELATED PROTEI RAP-1A; CHAIN: A; PROTO-ONKOGENE SERINE/THREONINE PROTEIN KINASE CHA B; TRANSFORMING PROT P21/H-RAS-1; CHAIN: A HIS-TAGGED TRANSFORMING PROT RANSFORMING PROT RHOA(0-181); CHAIN: A PKN; CHAIN: B; RAB6 GTPASE; CHAIN: ELONGATION PACTOR CHAIN: NULL;	RAS-RELATED PROTEI RAP-1A; CHAIN: A; PROTO-ONKOGENE SERINE/THREONINE PROTEIN KINASE CHA B; TRANSFORMING PROT P21/H-RAS-1; CHAIN: A HIS-TAGGED TRANSFORMING PROT RHOA(0-181); CHAIN: A PKN; CHAIN: B; RAB6 GTPASE; CHAIN: CHAIN: NULL; ADP-RIBOSYLATION FACTOR 6; CHAIN: A;
	60.17 TRANSFORMIN P21/H-RAS-1; CTRANSFORMIN P21/H-RAS-1; CTRANSFORMIN P21/H-RAS-1; CPKN; CHAIN: BRAB6 GTPASE; RAB6 GTPASE; CHAIN: NULL; CHAIN: NULL;	
TRANSFORMING PROTEIN RHOA(0-181); CHAIN: A; PKN; CHAIN: B; RAB6 GTPASE; CHAIN: A; ELONGATION FACTOR G; CHAIN: NULL;	TRANSFORMING PROTEIN RHOA(0-181); CHAIN: A; PKN; CHAIN: B; RAB6 GTPASE; CHAIN: A; ELONGATION FACTOR G; CHAIN: NULL;	TRANSFORMING PROTEIN RHOA(0-181); CHAIN: A; PKN; CHAIN: B; RAB6 GTPASE; CHAIN: A; ELONGATION FACTOR G; CHAIN: NULL; ADP-RIBOSYLATION FACTOR 6; CHAIN: A;
RAB6 GTPASE; CHAIN: A; ELONGATION FACTOR G; CHAIN: NULL;	ELONGATION FACTOR G;	ELONGATION FACTOR G; CHAIN: NULL; ADP-RIBOSYLATION FACTOR 6; CHAIN: A;
RAB6 GTPASE; CHAIN: A; ELONGATION FACTOR G; CHAIN: NULL;	RAB6 GTPASE; CHAIN: A; ELONGATION FACTOR G; CHAIN: NULL;	RAB6 GTPASE; CHAIN: A; ELONGATION FACTOR G; CHAIN: NULL; ADP-RIBOSYLATION FACTOR 6; CHAIN: A;
ELONGATION FACTOR (CHAIN: NULL;	ELONGATION FACTOR (CHAIN: NULL;	ELONGATION FACTOR C CHAIN: NULL; ADP-RIBOSYLATION FACTOR 6; CHAIN: A;
ELONGATION FACTOR CHAIN: NUILI;	ELONGATION FACTOR (CHAIN: NULL;	ELONGATION FACTOR (CHAIN: NULL; ADP-RIBOSYLATION FACTOR 6; CHAIN: A;
		ADP-RIBOSYLATION FACTOR 6; CHAIN: A;

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PDB annotation	TRAFFIC, GTP HYDROL YSIS, YPT/RAB 2 PROTEIN, ENDOCYTOSIS, HYDROL ASE	TRANSLATION EF-G; BENT CONFORMATION, VISIBLE DOMAIN III, MUTATION HIS573ALA	PROTEIN TRANSPORT GDP- BINDING, MEMBRANE TRAFFICKIN, NON- MYRISTOYLATED 1HUR 16	PROTEIN TRANSPORT GDP- BINDING, MEMBRANE TRAFFICKIN, NON- MYRISTOYLATED 1HUR 16	SMALL GTPASE KARYOPHERIN BETA, P95 SMALL GTPASE, NUCLEAR TRANSPORT RECEPTOR	GTP-BINDING PROTEIN, GTP-BINDING PROTEIN, SMALL G PROTEIN, RAP2, TOPP, RAS	GTP-BINDING PROTEIN, CTP-BINDING PROTEIN, SMALL G PROTEIN, RAP2, CDP, RAS	X JCLEAR STPASE, PORT	COMPLEX(GTPASE ACTIVATN/PROTO- ONCOGENE) GTPASE- ACTIVATING PROTEIN RHOGAP; COMPLEX
Compound		ELONGATION FACTOR G; CHAIN: A;	HUMAN ADP- RIBOSYLATION FACTOR 1; 1HUR 5 CHAIN: A, B; 1HUR 7	HUMAN ADP. RIBOSYLATION FACTOR 1; 1HUR 5 CHAIN: A, B; 1HUR 7	RAN; CHAIN: A, C; IMPORTIN BETA SUBUNIT; CHAIN: B, D;	RAP2A; CHAIN: NULL;	RAP2A; CHAIN: NULL;	RAN; CHAIN: A, C; NUCLEAR PORE COMPLEX PROTEIN NUP358; CHAIN: B, D;	PSO-RHOGAP; CHAIN: A; TRANSFORMING PROTEIN RHOA; CHAIN: B;
SEQ FOLD score				138.57	77.80	58.03		69.18	51.89
PMF score		0.03	1.00				1.00	·	
Verify score		-0.11	86.0				0.49		
Psi Blast		1.7e-05	le-64	1e-64	6.8e-48	3.40-51	3.4e-51	6.8e-48	le-34
END AA		137	185	185	186	.184	181	186	184
START AA		31	5	9	20	18	61	18	19
CHAIN		A	¥	¥	∢			ပ	æ
PDB U		Ifam	lhur	Ihur	libr	Ikao	Ikao	F1 F1	1tx4
SEQ ID NO:		371	371	371	371	371	371	371	371

Table 5

PDB annotation	(GTPASE ACTIVATION/PROTO- ONCOGENE), GTPASE, 2 TRANSITION STATE, GAP	COMPLEX (GTP-BINDING/EFFECTOR) RAS-RELATED PROTEIN RAB3A; COMPLEX (GTP-BINDING/EFFECTOR), GPROTEIN, EFFECTOR, RABCDR, 2 SYNAPTIC EXOCYTOSIS, RABPROTEIN, RAB3A, RABPHILIN	COMPLEX (GTP-BINDING/EFFECTOR) RAS-RELATED PROTEIN RAB3A; COMPLEX (GTP-BINDING/EFFECTOR, GPROTEIN, EFFECTOR, RABCDR, 2 SYNAPTICEXOCYTOSIS, RABPROTEIN, RAB3A, RABPHILIN	PROTEIN BINDING EF-G; EFF G ELONGATION FACTOR, TRANSLOCASE, RIBOSOME, ELONGATION, 2 TRANSLATION, PROTEIN SYNT FACTOR, GTPASE, GTP BINDING, 3 GUANOSINE NUCLEOTIDE BINDING, PROTEIN	HYDROLASE G PROTEIN, VESICULAR TRAFFICKING, GTP HYDROLYSIS, RAB 2 [1] PROTEIN, NEUROTRANSMITTER
Compound		RABPHILIN-3A; CHAIN: B;	RAB-3A; CHAIN: A; RABPHILIN-3A; CHAIN: B;	ELONGATION FACTOR G; CHAIN: A; ELONGATION FACTOR G DOMAIN 3; CHAIN: B;	RAB3A; CHAIN: A;
SEQ FOLD score		-	63.80		71.97
PMF score		1.00		0.35	
Verify		0.74		-0.05	
Psi Blast		1.7e-57	1.7e-57	3.4e-05	8.56-58
END		186	186	137	184
START		13	18	31	17
CHAIN		V.	Ą	A	A
PDB U			pqz1	2efg	3rab
SEQ ID NO:		371	371	371	371

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PDB annotation	RELEASE, HYDROLASE	HYDROLASE G PROTEIN,	VESICULAR TRAFFICKING,	GTP HYDROLYSIS, RAB 2	PROTEIN,	NEUROTRANSMITTER	KELEASE, HYDROLASE	HYDROL ASE	NET TO A MINITID A SEC.	NETRAMINIDASE	SIALIDASE, HYDROLASE	HYDROLASE	NEURAMINIDASE;	NEURAMINIDASE,	SIALIDASE, HIDROLASE	HYDROLASE	TRANSPORT A CE	GLYCOSIDASE	HYDROLASE	NEURAMINIDASE;	HYDROLASE,	GLYCUSIDASE	HYDROL ASE	NEURAMINIDASE;	HYDROLASE,	GLYCOSIDASE, SIGNAL, REPRAT. CALCIUM	HYDROLASE	NEURAMINIDASE;	HYDROLASE,	GLYCOSIDASE, SIGNAL,	HANDOI AGE	NEURAMINIDASE:	HYDROLASE	GLYCOSIDASE, SIGNAL,	100 Division 100 D
Compound		RAB3A; CHAIN: A;						STAT IDASE: CHAIN: NITLE				SIALIDASE; CHAIN: NULL;		-		SIALIDASE; CHAIN: NUIL;		-	SIALIDASE: CHAIN: NULL:				SIALIDASE; CHAIN: NULL;				SIALIDASE: CHAIN: NULL:				CTAT TO A SED. CUTAIN. NITT.	אבוטוגיים בישהויין ישנארודיוני			
SEQ FOLD score								135.41	-							99.82																			T
PMF		1.00			,							0.93							0.99				0.10				0.0-				5	C7:0			
Verify		0.72										0.48							0.46				0.43				0.13	}			900	86.0 0			
Psi Blast		8.5e-58			,			1 70-53	2			1.7e-53				0.00017			1.7e-53				0.0001				3.3e-17					3.46-12			
END AA		184						415	}			336				278			336				304		•		334	}			1	ş			
START AA		18						9	:			58				\$			58)			147				121	:			, 35	#			
CHAIN		Ą																																	1
PDB		3rab						1 Pull	3			1eur				leut			lent				1kit				1kit	į	-			ı Kat			
SEQ ID NO:		371						370	1	-		372				372			372	1			372				372	3				3/2			

Table 5

uo	OLASE,			PLEX), ZINC ING		PLEX), ZINC ING		PLEX), ZINC ing		JE	acres of		FINGER
PDB annotation	HYDROLASE HYDROLASE, INTRAMOLECULAR TRANS- SIALIDASE, NEURAMINIDASE	GLYCOSIDASE GLYCOSIDASE, HYDROLASE	GLYCOSIDASE GLYCOSIDASE, HYDROLASE	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN		COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN		COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN					COMPLEX (ZINC FINGER) PROTEIN-DNA
Compound	INTRAMOLECULAR TRANS-SIALIDASE; CHAIN: NULL;	SIALIDASE; CHAIN: NULL;	SIALIDASE; CHAIN: NULL;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B,	Ü	QGSR ZINC FINGER PEPTIDE, CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B,	<u>ن</u>	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B,	DNA-BINDING PROTEIN	HUMAN ENHANCER- BINDING PROTEIN MBP-1	MUTANT WITH CYS 11	1BBO 3 REPLACED BY ABU (C11ABU) (NMR, 60 STRICTIFIES: 1BBO 4	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;
SEQ FOLD score		92.46	·		·	65.40							
PMF score	0.05		0.03	0.25				0.35	40.0				0.18
Verify score	0:30		0.38	-0.32				0.35	-0.20		,		-0.27
Psi Blast	6.8e-32	8.5e-57	8.5e-57	3.4e-18		3.4e-26		3.4e-26	3.4e-07				1.7e-36
END	341	417	328	228		290		316	315				228
START AA	168	33	45	138	•	202		232	264				136
CHAIN				¥	-	∢		∢					ပ
PDB UI	2sli	3sil	3sil	lalh		laih		lalh	1bbo				Imey
SEQ ID NO:	372	372	372	373		373		373	373				373

PDB annotation	INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX	CZINC FUNGERDNA) COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX CZINC FINGER/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) TFIIIA; 5S GENE; NMR, TFIIIA, PROTEIN, DNA, TRANSCRIPTION FACTOR, 5S RNA 2 GENE, DNA BINDING PROTEIN, ZINC FINGER, COMPLEX 3 (TRANSCRIPTION	COMPLEX (TRANSCRIPTION) REGULATIONDNA) TFIIIA; [I] SS GENE; NMR, TFIIIA, [I] PROTEIN, DNA, TRANSCRIPTION FACTOR, (SS RNA 2 GENE, DNA BINDING PROTEIN, ZINC FINGER, COMPLEX 3 F	COMPLEX (TRANSCRIPTION)
Compound		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	TRANSCRIPTION FACTOR IIIA; CHAIN: A; 5S RNA GENE; CHAIN: B, F;	TRANSCRIPTION FACTOR IIIA; CHAIN: A; 58 RNA GENE; CHAIN: B, F;	TFIIIA; CHAIN: A, D; 5S
SEQ FOLD score		71.51		69.81		
PMF score			0.27		-0.01	0.00
Verify score			0.10		0.15	-0.33
Psi Blast		3.4e-41	3.4e-41	5.16-21	5.16-21	3.4e-24
END		289	316	292	315	236
START AA	·	201	231	201	232	101
CHAIN		ບ	U	4	· ·	A
PDB ID		Imey	Imey	£11.2	143	1tf6
SEQ ID NO:		373	373	373	373	373

PDB annotation	REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION) REGULATION/DNA) YING- YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA- PROTEIN RECOGNITION, 3 I
Сотроинд	RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	TFIIIA; CHAIN: A, D; SS RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	TFIILA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;
SEQ FOLD score				95.69	
PMF		0.04	0.70		0.24
Verify score		-0.09	0.04		-0.34
Psi Blast		1.5e-27	5.1e-38	5.1e-38	3.4e-26
AA END		267	315 ,	334	228
START AA		137	165	168	113
CHAIN		∢	V	∀	ပ
PDB U		1116	1466	146	lubd
SEQ ID NO:		373	373	373	373

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PDB annotation	REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION	INITIATION, INITIATOR FLEMENT, YY1, ZINC2	FINGER PROTEIN, DNA-	PROTEIN RECOGNITION, 3	COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA) YING- YANG 1: TRANSCRIPTION	INITIATION, INITIATOR	ELEMENT, YY1, ZINC 2	FINGER PROTEIN, DNA-	PROTEIN RECOGNITION, 3	COMPLEX (TRANSCRIPTION	REGULATION/DNA)	PECIT ATTOMORY VINC	YANG 1: TRANSCRIPTION	INITIATION, INITIATOR	ELEMENT, YYI, ZINC 2	FINGER PROTEIN, DNA-	PROTEIN RECOGNITION, 3	COMPLEX (TRANSCRIPTION)	REGULATION/DNA)	TRANSCRIPTION	REGULATION	TRANSCRIPTION	REGULATION, ADKI, ZINC I	COMPLEX (DNA-BINDING	PROTEIN/DIAN FIVE	ပ္	FINGER, COMPLEX (DNA- 17)	COMPLEX (DNA-BINDING	
Compound		YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT	DNA; CHAIN: A, B;	•			YY1; CHAIN: C; ADENO-	ASSOCIATED VIRUS P5	DNA; CHAIN: A, B;					Cimer O tarrio	A SSOCIATED AMERICA	ASSOCIATED VIKUS FO	DNA; CHAIN: A. B.						ADR1; CHAIN: NULL;				ZINC HINGER PROTEIN	GLII: CHAIN: A: DNA:	CHAIN: C, D;		ZINC FINGER PROTEIN	
SEQ FOLD score							89.17																				157 30					
PMF		0.98											,		0.0								.0.03								1.00	
Verify score		0.13						_						9	01.0								0.03								0.30	
Psi Blast		6.6e-34					1.2e-34							10.01	1.26-34		-						1.2e-10				6 Re-53	3		-	6.8e-53	
END		258					289							216	21.7								315				280	}			287	
START AA		4 1					176							700	200								264				136	}			168	
CHAIN		ບ					ပ							,	ر												\ \ \				V	
PDB III		1ubd					lubd							,	7001								2adr				2oli	i i			2gli	
SEQ ID NO:		373					373							717	6/6								373				373				373	

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			7			Z	acox' II	rsuc	- U-1	
PDB annotation	PROTEIN/DNA) FIVE- FINGER, GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE- FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	SERINE ESTERASE HYDROLASE, SERINE	ESTERASE, GLICOLEGICA	RNA-BINDING PROTEIN/RNA TRA PRE- MRNA; SPLICING REGULATION, RNP DOMAIN, RNA COMPLEX	GENE REGULATIONRNA POLY(A) BINDING PROTEIN 1, PABP 1; RRM, PROTEIN- RNA COMPLEX, GENE REGULATIONRNA	RNA BINDING PROTEIN RNA-BINDING DOMAIN	STRUCTURAL PROTEIN PROTEIN C23; RNP, RBD, RRM, RNA BINDING DOMAIN, NUCLEOLUS	STRUCTURAL PROTEIN PROTEIN C23; RNP, RBD, RRM, RNA BINDING DOMAIN, NUCLEOLUS	NUCLEAR PROTEIN HETEROGENEOUS NUCLEAR
Compound	GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	CUTINASE; CHAIN: NULL;		SXI-LETHAL PROTEIN; CHAIN: A, B; RNA (5'- R(P*GP*UP*UP*CP*UP*UP *UP*UP*UP*UP*UP*U)- CHAIN: P, Q;	POLYDENYLATE BINDING PROTEIN 1; CHAIN: A, B, C, D, E, F, G, H; RNA (5'- R(*AP*AP*AP*AP*AP*AP* AP*AP*AP*AP*A,3'); CHAIN: M, N, O, P, Q, R, S, T;	HU ANTIGEN C; CHAIN: A;	NUCLEOLIN RBD1; CHAIN: A;	NUCLEOLIN RBD2; CHAIN: A;	HNRNP A1; CHAIN: NULL;
SEQ FOLD score	·				·					. •
PMF		1.00	0.01		1.00	0.92	0.95	0.71	0.74	0.80
Verify score		0.27	-0.15		0.63	0.58	0.17	0.47	0.51	0.35
Psi Blast	•	3.4e-41	0.002		5.1e-13	3.4e-14	6.6e-14	1.3e-13	3,3e-14	1.7e-18
END AA		315	639		172	172	173	173	173	170
START AA	·	173	506		801	110	110	100	100	105
CHAIN		¥			¥	A	¥	А	V	
PDB U		2gli	lcex		167f	lcvj	1d8z	167	1fjc	1ha1
SEQ ID		373	374		376	376	376	376	376	376

					PLTA	502/0.		
PDB annotation	RIBONUCLEOPROTEIN A1, NUCLEAR PROTEIN, HNRNP, RBD, RRM, RNP, RNA BINDING, 2 RIBONUCLEOPROTEIN	RNA-BINDING PROTEIN RNA-BINDING DOMAIN		RNA-BINDING DOMAIN, RNA-BINDING DOMAIN, ALTERNATIVE SPLICING	COMPLEX (RIBONUCLEOPROTEIN/DN A) HNRNP A1, UP1; COMPLEX (RIBONUCLEOPROTEIN/DN A), HETEROGENEOUS NUCLEAR 2 RIBONUCLEOPROTEIN A1	RNA-BINDING PROTEINRNA TRA PRE- MRNA; SPLICING REGULATION, RNP DOMAIN, RNA COMPLEX		RNA-BINDING PROTEIN RNA-BINDING DOMAIN
Compound		HETEROGENEOUS NUCLEAR RIBONUCLEOPROTEIN DO; CHAIN: A;	RNA-BINDING PROTEIN SEX-LETHAL PROTEIN (C- TERMINUS, OR SECOND RNA-BINDING DOMAIN 1SXL 3 (RBD-2), RESIDUES 199 - 294 PLUS N- TERMINAL MET) 1SXL 4 (NMR, 17 STRUCTURES) 1SXL 5	SEX-LETHAL PROTEIN; CHAIN: NULL;	HETEROGENEOUS NUCLEAR RIBONUCLEOPROTEIN A1; CHAIN: A; 12- NUCLEOTIDE SINGLE- STRANDED TELOMETRIC DNA; CHAIN: B;	SXL-LETHAL PROTEIN; CHAIN: A, B; RNA (5'- R(P*GP*UP*UP*GP*UP *UP*UP*UP*UP*UP-U)- CHAIN: P, Q;	HU ANTIGEN C; CHAIN: A;	HU ANTIGEN C; CHAIN: A;
SEQ FOLD score							·	
PMF score		0.30	0.59	1.00	0.72	1.00	1.00	1.00
Verify		0.61	0.60	0.89	0.65	0.77	0.87	0.35
Psi Blast		8.5e-13	6.6e-13	5.1e-13	6.8e-20	5.1e-22	3.46-22	9.9e-25
AA END		166	173	172		192	192	193
START AA		111	105	801	104	110	107	110
CHAIN		Y			Α	A	4	A
PDB CI		1hd1	lsxl	2sxi	2up1	1b7f	1d8z	1d8z
SEQ ID NO:		376	376	376	376	377	377	377

Table 5

					TOTAL TOTAL MILES		
PDB annotation	RNA BINDING PROTEIN RNA-BINDING DOMAIN	STRUCTURAL PROTBIN PROTBIN C23; RNP, RBD, RRM, RNA BINDING	NUCLEAR PROTEIN HETEROGENEOUS NUCLEAR RIBONUCLEOPROTEIN, HURNP, RBD, RRM, RNP, RNA BINDING, 2 RIBONUCLEOPROTEIN	RNA BINDING PROTEIN RNA-BINDING DOMAIN			RNA-BINDING DOMAIN IL RNA-BINDING DOMAIN, IL ALTERNATIVE SPLICING FI
	RN/	STR PRO RRA	REAL PROPERTY OF THE PROPERTY	RN.		. `	RNA
Compound	HU ANTIGEN C; CHAIN: A;	NUCLEOLIN RBD1; CHAIN: A;	HNRNP A1; CHAIN: NUIL;	HETEROGENEOUS NUCLEAR RIBONUCLEOPROTEIN DO; CHAIN: A:	RNA-BINDING PROTEIN SEX-LETHAL PROTEIN (C- TERMINUS, OR SECOND RNA-BINDING DOMAIN 1SXL 3 (RBD-2), RESIDUES 199 - 294 PLUS N- TERMINAL MET) 1SXL 4 (NMR, 17 STRUCTURES) 1SXL 5	RNA-BINDING PROTEIN SEX-LETHAL PROTEIN (C- TERMINUS, OR SECOND RNA-BINDING DOMAIN ISXL 3 (RBD-2), RESIDUES 199 - 294 PLUS N- TERMINAL MBT) ISXL 4. (NMR, 17 STRUCTURES) ISXL 5	SEX-LETHAL PROTEIN; CHAIN: NULL;
SEQ FOLD score						51.72	57.11
PMF score	1.00	0.49	1.00	0.99	66'0		
Verify score	98.0	0.34	0.95	1.14	96'0		
Psi Blast	2.3e-22	9.9e-23	1.7e-30	1.4e-24	6.66-24	6.6e-24	5.1e-22
END AA	193	193	190	186	193	191	194
START AA	111	100	105	Ξ	105	. 86	107
CHAIN	A	ď		V		·	
EG e	1d9a	167	Ihal	1hd1	1sxl	Isxl	2sx1
SEQ ID NO:	27.1	377	377	377	377	377	377

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			_7 <i>5</i> :					18.75° 18.	the second second		TF 44	-
PDB annotation	RNA-BINDING DOMAIN RNA-BINDING DOMAIN, ALTERNATIVE SPLICING	COMPLEX (RIBONUCLEOPROTEIN/DN A) HNRNP A1, UP1;	(RIBONUCI EOPROTEIN/DN A), HETEROGENEOUS NUCI EAR 2 RIBONUCI EOPROTEIN A1	RNA BINDING DOMAIN RNA BINDING DOMAIN, RBD, RNA RECOGNITION MOTIF, RRM, 2 SPLICING	INHIBITOR, TRANSLATIONAL INHIBITOR, SEX 3	DETERMINATION, X CHROMOSOME DOSAGE COMPENSATION		PLANT PROTEIN TWO HOMOLOGOUS HEVEIN-	SUGAR BINDING PROTEIN " UDA; LECTIN, HEVEIN DOMAIN, UDA, SUPERANTIGEN	SUGAR BINDING PROTEIN (J. UDA; LECTIN, HEVEIN C. DOMAIN, UDA, SUPERANTIGEN, SACCHARIDE BINDING	SIGNALLING PROTEIN BINDING PROTEIN, CYTOKINE, SIGNALLING PROTEIN	COMPLEX
Compound	SEX-LETHAL PROTEIN; CHAIN: NULL;	HETEROGENEOUS NUCLEAR RIBONUCLEOPROTEIN A1;	NUCLEOTIDE SINGLE- STRANDED TELOMETRIC DNA; CHAIN: B;	SEX-LETHAL; CHAIN: A, B, C;		ï		AGGLUTININ ISOLECTIN VI; CHAIN: A	AGGLUTININ ISOLECTIN VI/AGGLUTININ ISOLECTIN V; CHAIN: A;	AGGLUTININ ISOLECTIN V/CHAIN: A;	TUMOR NECROSIS FACTOR RECEPTOR; CHAIN: A, B;	KALLIKREIN; CHAIN: A, B,
SEQ FOLD score				50.80								
PMF	1.00	1.00	·					0.21	0.48	0.45	90.0	0.24
Verify	. 06.0	1.01		·				-0.05	-0.30	0.38	-0.33	-0.56
Pst Blast	5.1e-22	6.8e-32		1.7e-21				1.7e-05	1.7e-05	1e-05	1.6e-07	0.00033
END	192	193	<u>-</u>	179				120	120	120	192	77
START AA	110	104		26				43	43	43	45	39
CHAIN		. V		∢				A	∢ .	· •	4	I
PDB U	2sxl	2up1		3sxl				1ehd	leis	len2	lext	1hia
SEQ ID NO:	377	377		377			! !	380	380	380	380	380

					_																		
PDB annotation	(PROTEASE/INHIBITOR) COMPLEX	(PROTEASE/INHIBITOR), TISSUE KALLIKREIN,	SERINE 2 PROTEASE,	TRYPSIN, PSA, KININ,	OK 1700 DE CHESTE	GLYCOPROTEIN	GLYCOPROTEIN GLYCOPROTEIN	GLYCOPROTEIN GLYCOPROTEIN	SERINE PROTEASE	INHIBITOR: ANTISTASIN,	CRYSTAL STRUCTURE,	HACTOR XA INHIBITOR, 2	NHIBITOR, THROMBOSIS				€ .eer	COMPLEX (GTPASE-	COMPLEX (GTPASE-	ACTIVATING/GTP-	ACTIVATION	• •	IRANSPORI, IRANSPORI R
Compound	X, Y; HIRUSTASIN; CHAIN: I, J;	·		•	THE STATE OF THE S	LAMININ; CHAIN: NULL;	LAMININ; CHAIN: NULL;	LAMININ; CHAIN: NULL;	ANTISTASIN; CHAIN:	NOLL;		•	-	LECTIN (AGGLUTININ) WHEAT GERM	AGGLUTININ (ISOLECTIN 2) 9WGA 3	LECTIN (AGGLUTININ) WHEAT GERM	AGGLUTININ (ISOLECTIN 2) 9WGA 3	P50-RHOGAP; CHAIN: A, B, C: CDC42HS; CHAIN: D, E.	Ä		•	GTP-BINDING PROTEIN RAN; CHAIN: A, B;	
SEQ FOLD score								73.03						96.99				66.17			•	97.94	
PMF .					8	-0.02	0.11		0.18							-0.19							
Verify score					9, 0	0.40	0.10		-0.21							-0.00							
Psi Blast					00 -3 0	8.56-09	3.3e-11	3.3e-11	1.7e-06					8.5e-09	,	8.5e-09		9.96-49				1.3e-58	
A END					97.	140	204	207	138					199		179		225				227	
START AA					,	3	43	43	39			•		41		7		31				29	
CHAIN														¥		¥_		Q				¥	
804 ID						1KIO	1klo	11410	lskz					в8м6		egw6		1am4				1byu	
SEQ ID NO:				-	700	380	380	380	380					380		380		385				385	

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		 1			121	The man is the	SUZ/	# # # # # # # # # # # # # # # # # # #	
PDB annotation	TRANSPORT PROTEIN TC4; GTPASE, NUCLEAR TRANSPORT, TRANSPORT PROTEIN	TRANSPORT PROTEIN TC4; GTPASE, NUCLEAR TRANSPORT, TRANSPORT PROTEIN	TRANSPORT PROTEIN TC4; GTPASE, NUCLEAR TRANSPORT, TRANSPORT PROTEIN	SIGNALING PROTEIN GTP- BINDING PROTEINS, PROTEIN-PROTEIN COMPLEX, EFFECTORS	SIGNALING PROTEIN GIP- BINDING PROTEINS, PROTEIN-PROTEIN COMPLEX, EFFECTORS	SIGNALING PROTEIN G PROTEIN, GTP HYDROLYSIS, KINETIC CRYSTALLOGRAPHY, 2 SIGNALING PROTEIN	ნ ე <u></u> ვ	SIGNALING PROTEIN PROTEIN-PROTEIN COMPLEX, ANTIPARALLEL [1] COLLED-COLL	ENDOCYTOSIS/EXOCYTOSI,
Compound	GTP-BINDING PROTEIN RAN; CHAIN: A, B;	GTP-BINDING PROTEIN RAN; CHAIN: A, B;	GTP-BINDING PROTEIN RAN; CHAIN: A, B;	RAS-RELATED PROTEIN RAP-1A; CHAIN: A; PROTO-ONKOGENE SERINE/THREONINE PROTEIN KINASE CHAIN: B;	RAS-RELATED PROTEIN RAP-1A; CHAIN: A; PROTO-ONKOGENE SERINE/THREONINE PROTEIN KINASE CHAIN: B;	TRANSFORMING PROTEIN P21/H-RAS-1; CHAIN: A;	TRANSFORMING PROTEIN P21/H-RAS-1; CHAIN: A;	HIS-TAGGED TRANSFORMING PROTEIN RHOA(0-181); CHAIN: A; PKN; CHAIN: B;	RAB6 GTPASE; CHAIN: A;
SEQ FOLD score		93.32		101.45		·	113.07	87.72	
PMF score	1.00		86'0		1.00	1.00			1.00
Verify	0.58		0.25		0.63	0.53			0.86
Psi Blast	1.3e-58	9.9e-59	9.96-59	3.4e-65	3.4e-65	3.4e-66	3,4e-66	6.8e-51	5.1e-59
END AA	211	227	211	201	201	201	202	201	199
START AA	32	26	27	31	32	33	33	28	34
CHAIN	Ą	æ	æ	Ą	¥	∀	Ą	V	V
PDB	1byu	1byu	1byu	lcly	lcly	lctq	lctg	lcxz	1d5c
SEQ ID NO:	385	385	385	385	385	385	385	385	385

Table 5

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PDB annotation	S G-PROTEIN, GTPASE, RAB6, VESICULAR TRAFFICKING	SIGNALING PROTEIN P21- RAC2; RHO GDI 2, RHO-GDI BETA IV GDI: BETA	SANDWHICH, PROTEIN- PROTEIN COMPLEX, G-	DOMAIN, 2	WALKER FOLD, GTP- BINDING PROTEIN	ENDOCYTOSIS/EXOCYTOSI	TRAFFIC, GTP	HYDROLYSIS, YPT/RAB 2	HYDROLASE	PROTEIN TRANSPORT GDP-	BINDING, MEMBRANE TRAFFICKTN NON-	MYRISTOYLATED 1HUR 16	SMALL GIPASE	SMALL GTPASE, NUCLEAR	TRANSPORT RECEPTOR	GTP-BINDING PROTEIN	GTP-BINDING PROTEIN,	GDP, RAS	GTP-BINDING PROTEIN GTP-BINDING PROTEIN	SMALL G PROTEIN, RAP2, TO GDP RAS	GTP-BINDING GTP-		G-PROTEIN, RHO FAMILY, MI RAS SUPER 2 FAMILY	A1
Compound		RAS-RELATED C3 BOTULINUM TOXIN STREETS ATTR 2: CHAIN: A:	RHO GDP-DISSOCIATION INHIBITOR 2: CHAIN: B:			GTP-BINDING PROTEIN YPT51: CHAIN: A:	fr : : : : : : : : : : : : : : : : : : :			HUMAN ADP.	KIBOSYLATION FACTOR	HUR 7	RAN; CHAIN: A, C;	SUBUNIT; CHAIN: B. D.		RAP2A; CHAIN: NULL;		•	RAP2A; CHAIN: NULL;		RACI; CHAIN: NUIL;			ONCOGENE PROTEIN C.H.
SEQ FOLD score										66.72			104.05			104.05					87.53			57.70
PMIF score		1.00			. '	1.00			- 1						. 1				1.00					
Verify		0.49				0.90	. ,				-								0.52					
Psi Blast		3.3e-53				le-56				1.7e-11			1.7e-51			1.7e-60			1.7e-60		1.2e-52			6.8e-51
END		204				199				203			207			202			199		211			201
START AA		32				34				21			33			31			32		29			33
CHAIN		Ą				A				A			Ą											
PDB ID		9sp1				1ek0				1hur			1ibr			1kao			lkao	***	ImhI			1plj
SEQ ID NO:		385				385				385			385			385			385		385	,		385

Table

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PDB annotation	·	COMPLEX (SMALL GIPASENUCLEAR PROTEIN) COMPLEX (SMALL GTPASENUCLEAR PROTEIN), SMALL GTPASE, 2 NUCLEAR TRANSPORT	COMPLEX(GTPASE ACTIVATN/PROTO- ONCOGENE) GTPASE- ACTIVATING PROTEIN RHOGAP; COMPLEX (GTPASE ACTIVATION/PROTO- ONCOGENE), GTPASE, 2 TRANSITION STATE, GAP	COMPLEX (GTP-BINDING/EFFECTOR) RAS- CAMPLEX (GTP-COMPLEX (GTP-BINDING/EFFECTOR), GPRABCTOR, CACCTOR, C	COMPLEX (GTP-BINDING/EFFECTOR) RAS- HARELATED PROTEIN RAB3AFIJ COMPLEX (GTP-BINDING/EFFECTOR), GRADING/EFFECTOR)
Compound	RAS P21 PROTEIN MUTANT WITH GLY 12 REPLACED BY PRO 1PLJ 3 (G12P) COMPLEXED WITH P3-1-(2- NITROPHENYL.)ETHYL- 1PLJ 4 GUANOSINE-5- (B,G-IMIDO)- TRIPHOSPHATE 1PLJ 5	RAN; CHAIN: A, C; NUCLEAR PORE COMPLEX PROTEIN NUP358; CHAIN: B, D;	P50-RHOGAP; CHAIN: A; TRANSFORMING PROTEIN RHOA; CHAIN: B;	RAB-3A; CHAIN: A; RABPHILIN-3A; CHAIN: B;	RAB-3A; CHAIN: A; RABPHILIN-3A; CHAIN: B;
SEQ FOLD score		104.86	73.28	115.76	
PMF score					1.00
Verify score				·	0.76
Psi Blast		8.5e-52	5.1e-48	1.7e-64	1.7e-64
END AA		221	199	203	206
START AA		31	31	23	30
CHAIN		ن د	Ø	A	A
PDB U		lтр	1tx4	1zbd	Izbd
SEQ ID NO:		385	385	385	385

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PDB annotation	PROTEIN, EFFECTOR, RABCDR, 2 SYNAPTIC EXOCYTOSIS, RAB PROTEIN, RAB3A,	HYDROLASE CDC42/CDC42GAP; CDC42/CDC42GAP; TRANSITION STATE, G- PROTEIN, GAP, CDC42, ALF3., HYDROLASE	HYDROLASE G PROTEIN, VESICULAR TRAFFICKÍNG, GTP HYDROLYSIS, RAB 2 PROTEIN, NEUROTRANSMITTER RELEASE, HYDROLASE	HYDROLASE G PROTEIN, VESICULAR TRAFFICKING, GTP HYDROLYSIS, RAB 2 PROTEIN, NEUROTRANSMITTER RELEASE, HYDROLASE	3	ENDOCYTOSIS/EXOCYTOSIJI S SYNAPTOTAGMIN ASSOCIATED 35 KDA PROTEIN, P35A, THREE HELIX BUNDLE	CONTRACTILE PROTEIN FOR TRIPLE-HELIX COLLED COLL, CONTRACTILE PROTEIN FOR THE
Compound		GTP BINDING PROTEIN (G25K); CHAIN: A; GTPASE ACTIVATING PROTEIN (RHG); CHAIN: B;	RAB3A; CHAIN: A;	RAB3A; CHAIN: A;	SYNTAXIN-1A; CHAIN: A, B, C;	SYNTAXIN-1A; CHAIN: A, B, C;	HUMAN SKELETAL MUSCLE ALPHA-ACTININ 2; CHAIN: A;
SEQ FOLD score	·	79.16		127.86			
PMF			1.00		-0.18	-0.17	-0.07
Verify			0.89		0.00	0.03	0.04
Psi Blast		9.9e-53	3.4e-65	3.4e-65	3.3e-08	2.3e-10	3.3e-18
END AA		229	202	202	292	227	292
START AA		31	29	30	195	94	21
CHAIN		V	4	V	A	¥	A
FOB EDB		2ngr	3rab	3rab	lez3	lez3	Iquu
SEQ ID NO:		385	385	385	386	386	386

	·			<u> </u>				
PDB annotation	COMPLEX (INHIBITOR/NUCLEASE) COMPLEX (INHIBITOR/NUCLEASE), COMPLEX (RI-ANG),	HYDROLASE 2 MOLECULAR RECOGNITION, EPITOPE MAPPING, LEUCINE-RICH 3 REPEATS	COMPLEX (INHIBITOR/NUCLEASE) COMPLEX (MHIBITOR/NUCLEASE)	COMPLEX (RI-ANG), HYDROLASE 2 MOLECULAR RECOGNITION, EPITOPE MAPPING, LEÜCINE-RICH 3 REPEATS	COMPLEX (INHIBITOR/NUCLEASE) COMPLEX	(INHIBITOR/NUCLBASE), COMPLEX (RI-ANG), HYDROLASE 2 MOLECULAR, RECOGNITION, EPITOPE MAPPING, LEUCINE-RICH 357 REPEATS	COMPLEX (NUCLEAR PROTEINRNA) COMPLEX (NUCLEAR PROTEINRNA), (I) RNA, SNRNP, RIBONUCLEOPROTEIN	COMPLEX (NUCLEAR PROTEIN/RNA) COMPLEX (NUCLEAR PROTEIN/RNA), HERNA, SNRNP, RIBONUCLEOPROTEIL SNRNP, RIBONUCLEOPROTEIL IN
Compound	RIBONUCLEASE INHIBITOR; CHAIN: A, D; ANGIOGENIN; CHAIN: B, E;		RIBONUCLEASE INFIBITOR; CHAIN: A, D; ANGIOGENIN; CHAIN: B, F.	î	RIBONUCLEASE INHIBITOR; CHAIN: A, D; ANGIOGENIN; CHAIN: B,	ŭ	U2 RNA HAIRPIN IV; CHAIN: Q, R; U2 A'; CHAIN: A, C; U2 B"; CHAIN: B, D;	UZ RNA HAIRPIN IV; CHAÎN: Q, R; U2 A; CHAÎN: A, C; U2 B"; CHAÎN: B, D;
SEQ FOLD score					·			
PIMF	0.37	· .	66.0		1.00		0.98	0.05
Verify score	-0.16		0.19		0.44		0.21	0.29
Psi Blast	8.5e-05		8.5e-14		2.6e-17		0.0037	0.0066
END AA	201		276		276		227	175
START AA	ជ		46		93		2 6	27
CHAIN	4		¥	·	¥		4	ပ
PDB U	la4y		la4y		la4y		1a9n	la9n
SEQ ID NO:	388		388		388		388	388

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PDB annotation	COMPLEX (NUCLEAR PROTEIN/RNA) COMPLEX (NUCLEAR PROTEIN/RNA), RNA, SNRNP, RIBONUCLEOPROTE	CELL ADHESION LEUCINE RICH REPEAT, CALCIUM BINDING, CELL ADHESION	CELL ADHESION LEUCINE RICH REPEAT, CALCIUM BINDING, CELL ADHESION	CONTRACTILE PROTEIN LEUCINE-RICH REPEAT,	BETA-BETA-ALPHA CYLINDER, DYNEIN, 2 CHLAMYDOMONAS,	FLAGELLA	CONTRACTILE PROTEIN LEUCINE-RICH REPEAT,	BETA-BETA-ALPHA	CHLAMYDOMONAS,	FLAGELLA	RNA BINDING PROTEIN TAP	RIBONUCL EOPROTEIN "	(RNP, RBD OR RRM) AND	(LRR)	RNA BINDING PROTEIN TAP	(NFAI);	(RNP, RBD OR RRM) AND	LEUCINE-RICH-REPEAT 2 GRR)	RNA BINDING PROTEIN TAM	(NFXI); RIBONUCLEOPROTEIN
Compound	UZ RNA HAIRPIN IV; CHAIN: Q, R; UZ A'; CHAIN: A, C; UZ B''; CHAIN: B, D;	INTERNALIN B; CHAIN: A;	INTERNALIN B; CHAIN: A;	OUTER ARM DYNEIN; CHAIN: A;			OUTER ARM DYNEIN; CHAIN: A;				NUCLEAR RNA EXPORT PACTOR 1: CHAIN: A R:		•		NUCLEAR RNA EXPORT	FACTOR 1; CHAIN: A, B;			NUCLEAR RNA EXPORT	FACTOR 1; CHAIN: A, B;
SEQ FOLD score		·									· .									
PMF	0.96	0.03	0.07	0.13			0.17				0.41				0.13				0.09	
Verify	60:0	-0.38	-0.22	0.07	·		0.17			,	0.05			į	0.12				-0.02	
Psi Blast	0.0037	8.5e-09	6.8e-14	16-07			le-07			9,	1.7e-07				1.7e-08				1.7e-08	
A EN	227	275	254	276			722				253				227				227	
START AA	42	150	88	135			<u>ಜ</u>				146				81				81	
CHAIN	υ_	¥.	¥	¥			∢				€				Ą				В	
PDB US	1a9n	140b	140b	6sp1	_	,	- Gsp I			,	lfo!				1fo1				Ifol	
SEQ ID NO:	388	388	388	388			388			000	388				388				388	

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PDB annotation	(RNP,RBD OR RRM) AND LEUCINE-RICH-REPEAT 2 (LRR)	LIGASE CYCLIN A/CDK2- ASSOCIATED PROTEIN P45; CYCLIN A/CDK2-	ASSOCIATED PROTEIN P19; SKP1, SKP2, F-BOX, 1.RR.	LEUCINE-RICH REPEAT,	SCF, UBIQUITIN, 2 E3, UBIQUITIN PROTEIN	LIGASE CYCLIN A/CDK2-	ASSOCIATED PROTEIN P45; CYCLIN A/CDK2-	ASSOCIATED PROTEIN P19;	SKP1, SKP2, F-BOX, LRR,	SCF. UBIOUITIN, 2 E3.	UBIQUITIN PROTEIN	LIGASE	LIGASE SKP2 F-BOX; SKP1; SKP1, SKP2, F-BOX, LRR,	LEUCINE-RICH REPEAT, "		LIGASE	ASSOCIATED P45: CYCLIN #	A/CDK2-ASSOCIATED P19;	SKPI, SKP2, F-BOX, LRRS,	LEUCINE-RICH REPEATS,	SCF, 2 UBIQUITIN, E3,	LIGASE	1000		SKP1 SKP2 R-ROY IRRS	J
Compound		SKP2; CHAIN! A, C, B, G, I, K, M, O; SKP1; CHAIN: B, D, F, H. J, L, N, P:				SKP2; CHAIN: A, C, E, G, I,	K, M, O; SKP1; CHAIN: B, D, F, H, J, L, N, P;				•		CYCLIN A/CDK2- ASSOCIATED P19; CHAIN:	A, C; CYCLIN A/CDK2-	B, D;	The state of the s	SKPZ; CHAIN: A, C; SKPI; CHAIN: B. D:						SKP2; CHAIN: A, C; SKP1;	CHAIN: B, D;		
SEQ FOLD score																										7
PMF score		1.00				0.24							0.70			,	3					,	0.39			
Verify score		0.25				0.28							-0.61				0.08						0.23			
Psi Blast		3.4e-33				1.7e-11							16-07			20	5.16-33						1.7e-11			
END		276				198							8	_		200	0/7						198			
START AA		25				∞							25			,	3						∞			<u> </u>
CHAIN		∢	-			A			-				∢				<						¥			
PDB ID		Ifqv				Ifqv							1fs1			0.3.	7911						1fs2			
SEQ ID NO:		388				388	·						. 388			000	388						388			1

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PDB annotation	LEUCINE-RICH REPEATS, SCF, 2 UBIQUITIN, E3, UBIQUITIN PROTEIN LIGASE	ACETYLATION RNASB INFIBITOR, PREOMITY PASSIANGIOGEN	N. INHIBITOR ACETYLATION, LEUCINE- RICH REPEATS	ACETYLATION RNASE INHIBITOR, PIBONICT BASE/ANGIOGEN	IN INHIBITOR ACETYLATION, LEUCINE- RICH REPEATS	ACETYLATION RNASE INHIBITOR,	RIBONUCLEASE/ANGIOGEN	ACETYLATION, LEUCINE- RICH REPEATS				COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC	FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX	(ZINC FINGERIDINA), ZINC FINGER, DNA-BINDING PROTEIN
Compound		RIBONUCLEASE INHIBITOR; CHAIN: NULL;		RIBONUCLEASE INHIBITOR; CHAIN: NULL;		RIBONUCI EASE INHIBITOR; CHAÎN: NULL;			LEUCINE ZIPPER GCN4 (BASIC REGION, LEUCINE	AP-1 DNA 1YSA 3		QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX	OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C:	OGSR ZINC FINGER PEPTIDE; CHAIN: A;	DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B,
SEQ FOLD score															
PMF score		0.62		0.71	•	0.95		-	-0.19			0.92		1.00	·
Verify		0.15		0.20	•	0.35			0.08			0.01		0.13	
Psi Blast		1.2e-12		1.4e-11		le-15			6.6e-16			5.1e <i>-27</i>	····	1.7e-29	
END		227	,	272		276			127			225		253	
START AA	·	13		2		46			73	·		155		173	
CHAIN									ပ			Ą		A	·
PDB ID		2bnh		2bnh		2bnh			lysa			iaih	· <u> </u>	laIh	
SEQ ID NO:	·	388		388		388			389			392		392	

Table 5

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PDB annotation		COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC	PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER) DNA) ZINC	HINGER, DNA-BINDING PROTEIN	COMPI RX (ZINC	FINGER/DNA) ZINC FINGER, PROTEIN-DNA	INTERACTION, PROTEIN DESIGN 2 CRYSTAL	STRUCTURE, COMPLEX	COMPLEX (ZINC	FINGER/DNA) ZINC FINGER, PROTEIN-DNA	INTERACTION, PROTEIN	DESIGN, 2 CRYSTAL	STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC BINGER L	PROTEIN-DNA	INTERACTION, PROTEIN	CENTRAL CRISIAL CONDINK	(ZINC FINGER/DNA)	COMPLEX (ZINC FINGER FI	PROTEIN-DNA	DESIGN, 2 CRYSTAL
Compound	C;	OGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX	OLIGONOCLEO I IDE BINDING SITE; CHAIN: B, C;	OGSR ZINC FINGER PEPTIDE; CHAIN: A;	OLIGONUCLEOTIDE BINDING SITE; CHAIN: B,	DNA: CHAIN: A B D F.	CONSENSUS ZINC FINGER PROTEIN; CHAIN: C. F. G.			DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER PROTEIN: CHAIN: C. P. G.				DNA; CHAIN: A, B, D, E;	PROTEIN; CHAIN: C, F, G;				DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;	
SEQ FOLD score		72.18													83.29							
PMF score				0.51		0 65	}			1.00										0.98		
Verify score				-0.07		02.0	3			0.13						-				0.11		
Psi Blast		1.7e-29	,	1.7e-24		3 40-41	•			1.7e-50	_				1.7e-50					1.26-40		
END AA		255		264		225				253					254					264		
START AA		173		201		154	·			172	_				172					200		
CHAIN		Ą		Ą		c)			Ü					၁					ပ		
80 E		lalh	•	lalh		1 mev				lmey					1mey			-		1mey		
SEQ ID NO:		392		392		300	7			392					392			-		392		

PDB annotation	STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER,	PROTEIN-DNA	DESIGN, 2 CRYSTAL	STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) TFIIIA;	SS GENE; NMR, TFIIIA,	PROTEIN, DNA,	S RNA 2 GENE DNA	BINDING PROTEIN, ZINC	FINGER, COMPLEX 3	(TRANSCRIPTION	REGULATION/DNA)	COMPLEX (TRANSCRIPTION	KEGULATION/DINA) IFILIA;	PROTEIN DNA.	TRANSCRIPTION FACTOR, THE	5S RNA 2 GENE, DNA	INC	FINGER, COMPLEX 3	(TRANSCRIPTION	COMPLEX (TRANSCRIPTION)	DECITE ATTONONA VIOLES	S GENE: NMR. TFIIIA.	PROTEIN, DNA.	N FACTOR,	S N	£3	REGULATION/DNA) FIL
Compound		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;	-			TRANSCRIPTION FACTOR IIIA; CHAIN: A; 5S RNA	GENE; CHAIN: E, F;			,				TRANSCRIPTION FACTOR	GENIE: CHAIN: A; 58 KNA	CENES, CITATIV. E. I.,	•					THE ANSCHEDE TON BACTOR	TINAL CHANGE A. SC DNA	GENE: CHAIN: E. F.					
SEQ FOLD score															56.76												•			
PMF		98.0	•				0.40																200	è:						
Verify		0.46					-0.01																500	20:0-						
Psi Blast		3.3e-09					6.8~16								1.26-19								1 25 10	1.40-19						
END		253					. 225								263								252	623	·	•				
START		228					162	_				. —			172								27.1	5/1						<u>.</u>
CHAIN		Ð					∢								∢								\ \	<						
PDB		1mey					E								143								Ę	<u> </u>						
SEQ ID		392					392								392		,						5	760						

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PDB annotation	COMPLEX (TRANSCRIPTION REGULATION/DNA) THILA; SS GENE; NMR, THILA, PROTEIN, DNA, TRANSCRIPTION PACTOR	5S RNA 2 GENE, DNA BINDING PROTEIN, ZINC FINGER, COMPLEX 3 (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION TRANSCRIPTION INTIATION, ZINC FINGER	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATOR ELMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOMITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION* REGULATION/DNA) YING- [] YANG 1; TRANSCRIPTION [] INITIATION, INITIATOR []
Compound	TRANSCRIPTION FACTOR IIIA; CHAIN: A; SS RNA GENE; CHAIN: E, F;		TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;
SEQ FOLD score				53.26	66.82	·
PMF score	0.43		0.01			0.98
Verify score	-0.12	·	-0.39			90:0-
Psi Blast	1.70-17	·	1.7e-27	1.7e-27	6.86-31	6.8e-31
AA ES	266		262	757	254	253
START AA	201		162	72	140	156
CHAIN	⋖		4	.	د	ပ
EGE EGE	E		1466	1116	1ubd	lubd
SEQ ID NO:	392		392	392	392	392

Table 5

				P	OT/CSO	2/012	
PDB annotation	ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA- PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRETION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING- YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA- PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE- FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE- FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	MPLEX NA), ZINC NDING	MPLEX NA), ZINC NDING	COMPLEX (ZINC FINGER/DNA) COMPLEX
Compound		YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	OGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A;
SEQ FOLD score				58.63			
PMF		0.60	0.94		-0.20	-0.11	0.21
Verify		-0.04	-0.16		0.37	0.42	0.35
Psi Blast		1.7e-28	5.16-29	5.1e-29	le-17	5.1e-22	6.8e-27
END A		261	252	254	747	781	608
START	·	180	152	66	699	202	723
CHAIN		υ	. •	∀	V	V	A
PDB		lubd	2gli	2gli	lalh	laih	lalh
SEQ ID		332	392	392	393	393	393

Table

PDB annotation	(ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN				LIGASE CBL, UBCH7, ZAP- 70, E2, UBIQUITIN, E3, PHOSPHORYLATION, 2 TYROSINE KINASE, UBIQUITINATION, PROTEIN	METAL BINDING PROTEIN NEING FINGER PROTEIN COMATI; RING FINGER COHC4)	COMPLEX (ZINC IV FINGER I I PROTEIN-DNA II II III III III III III IIIIIIIIII
Compound	DUPLEX OLIGONUCLEOTIDE BINDING SITE, CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	DNA-BINDING PROTEIN HUMAN ENHANCER- BINDING PROTEIN MBP-1 MUTANT WITH CYS 11 1BBO 3 REPLACED BY ABU (C11ABU) (NMR, 60 STRUCTURES) 1BBO 4	VIRUS EQUINB HERPES VIRUS-1 (C3HC4, OR RING DOMAIN) 1CHC 3 (NMR, 1 STRUCTURE) 1CHC 4	VIRUS EQUINE HERPES VIRUS-1 (C3HC4, OR RING DOMAIN) 1CHC 3 (NMR, 1 STRUCTURE) 1CHC 4	SIGNAL TRANSDUCTION PROTEIN CBL; CHAIN: A; ZAP-70 PEPTIDE; CHAIN: B; UBIQUITIN- CONTUGATING ENZYME E12-18 KDA UBCH7; CHAIN: C;	CDK-ACTIVATING KINASE ASSEMBLY FACTOR MAT1; CHAIN: A;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;
SEQ FOLD : score			·					·
PMF score		0.78	0.51	0.25	0.11	0.18	0.18	-0.13
Verify score		0.33	-0.28	-0.77	-0.44	-0.29	-0.13	0.29
Psi Blast		8.5e-28	2e-14	0.00013	0.0097	0.0097	6.6e-05	3.4e-40
END		833	813	35	56	22	63	781
START AA		757	759	11	11	11	11	695
CHAIN		A				¥	∢	ນ
BOY CI		lalh	1bbo	1chc	1chc	1fbv	1825	Ітеу
SEQ ID NO:		393	393	393	393	393	393	393

Table

PDB annotation	INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	DNA-BINDING PROTEIN V(D)J RECOMBINATION ACTIVATING PROTEIN 1; RAG1, V(D)J RECOMBINATION, ANTIBODY, MAD, RING FINGER, 2 ZINC BINUCLEAR CLUSTER, ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (TRANSCRIPTION REGULATION SS GENE; NMR, TFIIIA, [1] PROTEIN, DNA, [1] TRANSCRIPTION FACTOR, [1]
Compound		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	RAGI; CHAIN: NULL;	TRANSCRIPTION FACTOR IIIA; CHAIN: A; 58 RNA GENE; CHAIN: E, F;
SEQ FOLD score						
PMF score		0.64	0.99	0.95	0.33	0.62
Verify score		0.37	0.58	0.08	-0.66	0.29
Psi Blast		5.1e-49	5.1e-50	1.5e-13	0.00051	3.4e-19
END		608	835	608	35	833
START		22L	756	782	ო	757
CHAIN		ບ	U	₀		V
PDB	 	1mey	Imey	lmey	1rmd	£ .
SEQ ID		393	393	393	393	393

Table 5

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PDB annotation	SS RNA 2 GENE, DNA BINDING PROTEIN, ZINC FINGER, COMPLEX 3	(TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION	POLYMERASE III, 2	TRANSCRIPTION	INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION	REGULATION/DNA) COMPLEX (TRANSCRIPTION	REGULATION/DNA), RNA	POLYMERASE III, 2	TRANSCRIPTION	INITIATION, ZINC FINGER	NOTITE ANSCRIPTION I	REGULATION/DNA)	COMPLEX (TRANSCRIPTION)	REGULATION/DNA), RNA	POLYMERASE III, 2	TRANSCRIPTION	INTITATION, ZINC FINGER	COMPLEX (TRANSCRIPTION)	REGULATION/DNA)	COMPLEX (TRANSCRIPTION)	REGULATION/DNA), KNA	TO ANGUETION	TAITTATION ZINC BINGER	PROTEIN	COMPLEX (TRANSCRIPTION) REGULATION/DINA) YING- FI
Compound			TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE;	CHAIN: B, C, E, F;		•		TFIIIA; CHAIN: A, D; 5S	RIBOSOMAL RNA GENE; CHAIN: B. C. E. F.			•		THERE'S CHAIN. A D. CC.	RIBOSOMAL RNA GENE:	CHAIN: B, C, E, F;					TFIIIA; CHAIN: A, D; 5S	RIBOSOMAL RNA GENE;	CHAIN: B, C, E, F;					YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS P5
SEQ FOLD score																												
PMF score			-0.20					-0.19						27.0) 						-0.15							-0.20
Verify score			0.14					0.18						8	3						0.22							0.07
Psi Blast			5.1e-13					5.1e-21				•		2 42.20	3.46-23			•			3.4e-26							1.7e-19
END AA			762					790						010	010						835							748
START AA			647					657			•			077	900					J	723							647
CHAIN			¥	- -				Ą						_	<						A				٠.			ပ
PDB UD			1476					1tf6						1466	omi						1466							lubd
SEQ ID NO:			393					393						202	252						393							393

Table 5

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PDB annotation	YANG 1; TRANSCRIPTION INTTATION, INTTATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA- PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING- YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA- PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIFTION REGULATION/DNA) YING-YANG 1; TRANSCRIFTION INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION) REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 [1] COMPLEX (TRANSCRIPTION, REGULATION/DNA)	TRANSCRIPTION REGULATION TRANSCRIPTION REGULATION, ADRI, ZINC [1] FINGER, NMR
Compound	INITIATOR BLEMENT DNA; CHAIN: A, B;	YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DINA; CHAIN: A, B;	YYI; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 INTTATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	ADRI; CHAIN: NULL;
SEQ FOLD score	-				
PMF score		-0.17	-0.01	0.22	0.76
Verify score		0.28	0.33	0.37	0.30
Psi Blast		16-23	1e-29	3.46-32	5.1e-16
END		781	608	833	811
START AA		675	869	730	757
CHAIN ID		v	U	U	
PDB ID		1ubd	lubd	lubd	2adr
SEQ ID NO:		393	393	393	393

PDB annotation	TRANSCRIPTION REGULATION TRANSCRIPTION REGULATION, ADRI, ZINC FINGER, NMR		COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE. FINGER GIL; GIL, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE- FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GLJ; GLJ, ZINC FINGER, COMPLEX (DNA-FINGER, COMPLEX (DNA-FINDING) FINDING PROTEIN/DNA)	GENE REGULATION POZ DOMAIN; PROTEIN- PROTEIN INTERACTION DOMAIN, TRANSCRIPTIONAL 2 REPRESSOR, ZINC-FINGER PROTEIN, X-RAY CRYSTALLOGRAPHY, 3 PROTEIN STRUCTURE, PROMYELOCYTIC LEUKEMIA, GENE
Compound	ADR1; CHAIN: NULL;	COMPLEX(TRANSCRIPTIO N REGULATION/DNA) TRAMTRACK PROTEIN (TWO ZINC-FINGER PEPTIDE) COMPLEXED WITH 2DRP 3 DNA 2DRP 4	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLI1; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	PROMYBLOCYTIC LEUKEMIA ZINC FINGER PROTEIN PLZF; CHAIN: A;
SEQ FOLD score				·		
PMF score	0.11	0.94	-0.20	-0.12	0.30	1.00
Verify	-0.28	0.33	0.25	0.12	0.29	0.22
Psi Blast	8.5e-14	9.9e-16	1.2e-22	1.7e-27	1.2e-30	3.4e-22
END A	833	811	780	808	833	141
START	787	757	650	672	702	18
CHAIN		V	Y	V	· V	∢
PDB UD	2adr	2drp	2gli	2gli	2gli	1buo
SEQ ID NO:	393	393	393	393	393	394

Cable :

					Africa	~	
PDB annotation	GENE REGULATION POZ DOMAIN; PROTEIN- PROTEIN INTERACTION DOMAIN, TRANSCRIPTIONAL 2 REPRESSOR, ZINC-FINGER PROTEIN, X-RAY CRYSTALLOGRAPHY, 3 PROTEIN STRUCTURE, PROMYBLOCYTIC LEUKEMIA, GENE REGULATION		ENDOCYTOSIS/EXOCYTOSI S SYNAPTOTAGMIN, C2- DOMAIN, EXOCYTOSIS, NEUROTRANSMITTER 2 RELEASE, ENDOCYTOSIS/EXOCYTOSI	ENDOCYTOSIS/EXOCYTOSI (S SYNAPTOTAGMIN, C2- DOMAIN, EXOCYTOSIS, NEUROTRANSMITTER 2 RELEASE, ENDOCYTOSIS/EXOCYTOSI	ENDOCYTOSIS/EXOCYTOSI [1] S BETA SANDWICH,	ENDOCYTOSIS/EXOCYTOSI S S BETA SANDWICH, CALCIUM ION, C2 DOMAIN II	TRANSFERASE CALCIUM++
Compound	PROMYBLOCYTIC LEUKEMIA ZINC FINGER PROTEIN PLZF; CHAIN: A;	OXIDOREDUCTASE(OXYG EN(A)) GALACTOSE OXIDASE (B.C.1.1.3.9) (PH 4.5) 1GOF 3	SYNAPTOTAGMIN I; CHAIN: A;	SYNAPTOTAGMIN I; CHAIN: A;	SYNAPTOTAGMIN III; CHAIN: A;	SYNAPTOTAGMIN III; CHAIN: A;	PROTEIN KINASE C, ALPHA TYPE; CHAIN: A;
SEQ FOLD score	73.07						
PMF score		0.76	1.00	0.1	1.00	1.00	0.36
Verify score		0.13	0.88	0.74	09.0	0.48	0.19
Psi Blast	3,46-22	le-17	1.36-40	3.4e-34	1.7e-87	1.5e-57	le-20
END	144	587	274	269	418	417	427
START AA		306	141	146	141	146	296
CHAIN	¥		4	∢	4	4	A
PDB ID	1buo	lgof	1byn	1byn	ldqv	ldqv	ldsy
SEQ ID NO:	394	394	395	395	395	395	395

Table 5

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PDB annotation	PROTBIN, CALCIUM- BINDING 2-PROTBIN, PHOSPHATIDYLSERINE, PROTBIN KINASE C				ANTI-ONCOGENE CELL CYCLE, ANTI-ONCOGENE, REPEAT, ANK REPEAT	COMPLEX (TRANSCRIPTION REGULATION DNA) GABPALPHA; GABPBETAI; COMPLEX (TRANSCRIPTION REGULATION DNA-BINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, ANKYRIN REPEATS, TRANSCRIPTION 3 PACTOR I	COMPLEX (TRANSCRIPTION: REGULATION/DNA) GABPALPHA; GABPBETA1; L COMPLEX (TRANSCRIPTION; REGULATION/DNA), DNA- BINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, [1]
Compound		CALCIUM/PHOSPHOLIPID BINDING PROTEIN SYNAPTOTAGMIN I (FIRST C2 DOMAIN) (CALB) IRSY 3	CALCIUM/PHOSPHOLIPID BINDING PROTEIN SYNAPTOTAGMIN I (FIRST C2 DOMAIN) (CALB) IRSY 3	CALCTUM/PHOSPHOLIPID BINDING PROTEIN SYNAPTOTAGMIN I (FIRST C2 DOMAIN) (CALB) IRSY 3	TUMOR SUPPRESSOR P16INK4A; CHAIN: NULL;	GA BINDING PROTEIN ALPHA; CHAIN: A; GA BINDING PROTEIN BETA 1; CHAIN: B; DNA; CHAIN: D, E;	GA BINDING PROTEIN ALPHA; CHAIN: A; GA BINDING PROTEIN BETA 1; CHAIN: B; DNA; CHAIN: D, E;
SEQ FOLD score			169.67				·
PMF score		1.00		1.00	1.00	0.03	1.00
Verify score		0.70		62.0	0.20	-0.10	-0.13
Psi Blast		9.9e-46	9.9e-46	3.4e-34	6.6e-23	5.1e-34	1.2e-33
END		271	272	269	124	323	224
START AA		132	132	146	. 10	180	52
CHAIN	·	·				Ф	Я
PDB ID		Irsy	Irsy	Irsy	la5e	lawc	Iawc
SEQ ID NO:		395	395	395	396	396	396

	×	NO		NO		-	~	 ജ			•			-	7							Barre			s II	le «	<u> </u>	ALP ED	Ψ.			E	7
PDB annotation	EATS, ON 3 FACT(ANSCRIPTI	GABPBETA	ANSCRIPTI	DNA), DNA	CLEAR	DOMAIN,	EATS, ON 3 FACTO	ESSOR	ESSOR,	I S.	ESSOR	ESSOR,	TOR,	TE.	ESSOR	TOR.		VASE/ANTI	DK6;	SI; CYCLIN	INASE,	IDENT.	TORY 2	, INK4, CEL	PLE TUMO	MILSI,	HASE/ANTI-	TRITOR	SE)	OTEIN,	(DENT	4 1 1 1
PDB an	ANKYRIN REPEATS, TRANSCRIPTION 3 FACTOR	COMPLEX (TRANSCRIPTION REGIT ATTON/ONA)	GABPALPHA; GABPBETA1;	COMPLEX (TRANSCRIPTION	REGULATION/DNA), DNA-	BINDING, 2 NUCLEAR	FROIEIN, EIS DOMAIN	ANK YRIN REPEATS, TRANSCRIPTION 3 FACTOR	TUMOR SUPPRESSOR	TUMOR SUPPRESSOR,	ANKYRIN MOTIF	TUMOR SUPPRESSOR	TUMOR SUPPRESSOR,	CDK4/6 INHIBITOR,	ANKYRIN-MOTIF	TUMOR SUPPRESSOR	CDK4/6 INHIBITOR.	ANKYRIN MOTIF	COMPLEX (KINASE/ANTI-	ONCOGENE) CDK6;	P16INK4A, MTS1; CYCLIN	DEPENDENT KINASE,	CYCLIN DEPENDENT	KINASE INHIBITIORY 2	PROTEIN, CDK, INK4, CELL	CYCLE, MULTIPLE TUMOR	JEFRESSOR,	COMPLEX (KINASE/ANTI- ONCOGENE) HEADED	COMPLEX (INHIBITION	PROTEIN/KINASE)	INHIBITOR PROTEIN	CYCLIN-DEPENDENT	יייייי (יייייי
-	AE	O P			24 1	<u> </u>	<u> </u>	4 E	H			-		<u></u>	₹			A	D 	<u></u>	<u></u>		<u>ပါ</u>	Z	<u> </u>	<u>್</u>	ਰ ਹੋ -	<u>ა</u> ნ) E		Z	6 5	
Compound		GA BINDING PROTEIN AI PHA: CHAIN: A: GA	BINDING PROTEIN BETA	I; CHAIN: B; DNA; CHAIN:				:) CDK4/6	INHIBITOR; CHAIN: NULL;) CDK4/6	INHIBITOR; CHAIN: NULL;			PISINKAD CDK4/6 INHIBITOR: CHAIN: NIII I :			CYCLIN-DEPENDENT	KINASE 6; CHAIN: A;	MULTIPLE TUMOR	SUPPRESSOR; CHAIN: B;			٠.				CYCL IN-DEPENDENT	KINASE 6; CHAIN: A;	P19INK4D; CHAIN: B;		
		GA BIND ALPHA:	BINDING	1; CHAIN	ر بر بر		•		P19INK4D CDK4/6	INHIBITIC		P19INK4D CDK4/6	NHIBITO			PISINKAD CDK4/6 NHIBITOR: CHAP			CYCLIN-	KIINASE (MULTIPL	SUPPRES			•				CYCL IN-1	KINASE 6	P19INK4L		
SEQ FOLD score								-					į.											.•									
PMF score		1.00.					·		0.88			1.00				18.0			1.00										1.00				
Verify score		0.16							0.14			60.0				0.21	-		0.25										0.28	ļ.			
Psi Blast		6.8e-35							1.3e-30			le-27			,	9.96-31			1.6e-20						_				1.2e-26				
END AA		195		•					198			195			200	077			102										195		_		
START AA		6							10			12			ę	9			9	-									12				
CHAIN		В																	m										В				
908 E1		lawc							1pq8			1bd8			11,40	onat	_		1bi7										1blx				
SEQ ID NO:		396							396			396		-	306	060	•		396				•						396				

Table 5

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PDB annotation	CONTROL, ALPHA/BETA, COMPLEX (INHIBITOR PROTEIN/KINASE)	COMPLEX (INHIBITOR PROTEIN/KINASE) INHIBITOR PROTEIN, CYCLIN-DEPENDENT KINASE, CELL CYCLE 2 CONTROL, ALPHA/BETA, COMPLEX (INHIBITOR PROTEIN/KINASE)	COMPLEX (INHIBITOR PROTEIN/KINASE) INHIBITOR PROTEIN, CYCLIN-DEPENDENT KINASE, CELL CYCLE 2 CONTROL, ALPHA/BETA, COMPLEX (INHIBITOR PROTEIN/KINASE)	HORMONE/GROWTH FACTOR P18-INK4C; CELL CYCLE INHIBITOR, P18INK4C, TUMOR, SUPPRESSOR, CYCLIN-2 DEPENDENT KINASE, HORMONE/GROWTH FACTOR	CELL CYCLE INHIBITOR P18-INK4C(INK6); CELL CYCLE INHIBITOR, P18- INK4C(INK6), ANKYRIN REPEAT, 2 CDK 4/6 INHIBITOR	CELL CYCLE INHIBITOR P18-INK4C(INK6); CELL CYCLE INHIBITOR, P18- INK4C(INK6), ANKYRIN REPEAT, 2 CDK 4/6 INHIBITOR
Compound		CYCLIN-DEPENDENT KINASE 6, CHAIN: A; PI9INK4D; CHAIN: B;	CYCLIN-DEPENDENT KINASE 6; CHAIN: A; P19INK4D; CHAIN: B;	CYCLIN-DEPENDENT KINASE 6 INHIBITOR; CHAIN: A;	CYCLIN-DEPENDENT KINASE 6 INHIBITOR; CHAIN: A, B;	CYCLIN-DEPENDENT KINASE 6 INHIBITOR; CHAIN: A, B;
SEQ FOLD score						
PMF score		0.21	1.00	0.39	-0.17	1.00
Verify score		0.06	0.29	0.08	90:0	0.27
Psi Blast	-	3e-28	9.9e-34	1.7e-32	1.5e-28	16-31
END		226	202	200	322	199
START AA		50	رم ا	6	183	6
CHAIN ID	-	œ.	я	V	A	A
PDB ID		1blx	1blx	1bu9	libb	lihb
SEQ ID NO:		396	396	396	396	396

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PDB annotation	TRANSCRIPTION FACTOR P65; P50D; TRANSCRIPTION FACTOR, IKB/NFKB COMPLEX	ANK-REPEAT MYOTROPHIN, ACETYLATION, NMR, ANK- REPEAT	ANK-REPBAT MYOTROPHIN, ACETYLATION, NMR, ANK- REPEAT	COMPLEX (TRANSCRIPTION REG/ANK REPEAT) COMPLEX (TRANSCRIPTION REGULATION/ANK REPEAT), ANKYRIN 2 REPEAT HELIX	TRANSCRIPTION REGULATION TRANSCRIPTION REGULATION, ANKYRIN REPEATS, CELL, CYCLE	LB CR,	COMPLEX (TRANSCRIPTION)
Compound	NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF- KAPPA-B P50D SUBUNIT; CHAIN: C; I-KAPPA-B- ALPHA; CHAIN: D;	MYOTROPHIN; CHAIN: NULL	MYOTROPHIN; CHAIN: NULL	NF-KAPPA-B P65; CHAIN: A, C; NF-KAPPA-B P50; CHAIN: B, D; I-KAPPA-B- ALPHA; CHAIN: B, F;	REGULATORY PROTEIN SWI6; CHAIN: A, B;	P53; CHAIN: A; 53BP2; CHAIN: B;	GA BINDING PROTEIN
SEQ FOLD score							66.19
PMF score	0.94	1.00	0.53	0.77	0.99	0,54	
Verify	-0.06	-0.02	-0.16	-0.05	-0.08	-0.27	
Psi Blast	6.8e-38	1.4e-23	1.2e-22	3.4e-39	2e-32	1.3e-25	8.5e-39
END AA	235	135	164	235	218	217	152
START AA	4	10	53		-	σ.	16
CHAIN	D			m	V	m	В
PDB ID	likn	Ітуо	Imyo	lafi	1sw6	1 ycs	lawc
SEQ ID NO:	396	396	396	396	396	396	397

Table :

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PDB annotation	REGULATION/DNA) GABPALPHA; GABPBETA1; COMPLEX (TRANSCRIPTION REGULATION/DNA), DNA- BINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, ANKYRIN REPEATS, TRANSCRIPTION 3 HACTOR	COMPLEX (TRANSCRIPTION REGULATION/DNA) GABPALPHA; GABPBETAI; COMPLEX (TRANSCRIPTION REGULATION/DNA), DNABINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, ANKYRIN REPEATS, TRANSCRIPTION 3 FACTOR	COMPLEX (TRANSCRIPTION REGULATION/DNA) GABPALPHA; GABPBETA1; COMPLEX (TRANSCRIPTION REGULATION/DNA), DNABINDING, 2 NUCLEAR PROTEIN, ETS DOMAIN, TANSCRIPTION 3 PACTOR	TUMOR SUPPRESSOR, TUMOR SUPPRESSOR, CDK4/6 INHIBITOR, ANKYRIN MOTIF	COMPLEX (INHIBITOR PROTEIN/KINASE) INHIBITOR PROTEIN, CYCLIN-DEPENDENT KINASE, CELL CYCLE 2 CONTROL, ALPHA/BETA, COMPLEX (INHIBITOR PROTEIN/KINASE)	HORMONE/GROWTH
Compound	ALPHA; CHAIN: A; GA BINDING PROTEIN BETA 1; CHAIN: B; DNA; CHAIN: D, E;	GA BINDING PROTEIN ALPHA; CHAIN: A; GA BINDING PROTEIN BETA 1; CHAIN: B; DNA; CHAIN: D, E;	GA BINDING PROTEIN ALPHA; CHAIN: A; GA BINDING PROTEIN BETA 1; CHAIN: B; DNA; CHAIN: D, E;	P19INK4D CDK4/6 INHIBITOR; CHAIN: NULL;	CYCLIN-DEPENDENT KINASE 6; CHAIN: A; P19INKAD; CHAIN: B;	CYCLIN-DEPENDENT
SEQ FOLD score						56.15
PMF score		1.00	1.00	1.00	1.00	
Verify score		9.64	99.0	0.55	0.38	
Psi Blast		8.56-39	6.8e-37	1.5e-31	5.1e-30	1.2e-35
END		136		139	139	149
START AA		-	22	1		-
CHAIN		æ			а	A
PDB ID		lawc	lawc	1bd8	16kx	1bu9
SEQ ID NO:		397	397	397	397	397

Table 5

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PDB annotation	FACTOR P18-INK4C; CELL CYCLE INHIBITOR, P18INK4C, TUMOR, SUPPRESSOR, CYCLIN-2 DEPENDENT KINASE, HORMONE/GROWTH FACTOR	HORMONE/GROWTH PACTOR P18-INK4C; CELL CYCLE INHIBITOR, P18INK4C, TUMOR, SUPPRESSOR, CYCLIN- 2 DEPENDENT KINASE, HORMONE/GROWTH PACTOR	CELL CYCLE INHIBITOR P18-INK4C(INK6); CELL CYCLE INHIBITOR, P18- INK4C(INK6), ANKYRIN REPEAT, 2 CDK 4/6 INHIBITOR	CELL CYCLE INHIBITOR P18-INK4C(INK6); CELL CYCLE INHIBITOR, P18- INK4C(INK6), ANKYRIN REPEAT, 2 CDK 4/6 INHIBITOR	TRANSCRIPTION FACTOR P65; P50D; TRANSCRIPTION FACTOR, IXB/NFKB COMPLEX	TRANSCRIPTION FACTOR Y P65; P50D; TRANSCRIPTION F FACTOR, IKB/NFKB COMPLEX	ANK-REPEAT MYOTROPHIN,
Compound	KINASE 6 INFIBITOR; CHAIN: A;	CYCLIN-DEPENDENT KINASE 6 INHIBITOR; CHAIN: A;	CYCLIN-DEPENDENT KINASE 6 INHIBITOR; CHAIN: A, B;	CYCLIN-DEPENDENT KINASE 6 INHIBITOR; CHAIN: A, B;	NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF- KAPPA-B P50D SUBUNIT; CHAIN: C; I-KAPPA-B- ALPHA; CHAIN: D;	NF-KAPPA-B P65 SUBUNIT; CHAIN: A; NF- KAPPA-B P50D SUBUNIT; CHAIN: C; I-KAPPA-B- ALPHA; CHAIN: D;	MYOTROPHIN; CHAIN: NULL
SEQ FOLD score		·	56.08				52.07
PMF		00:1		1.00	1.00	0.81	
Verify score		0.40		0.51	0.37	0.21	
Psi Blast		1.2e-35	5.1e-35	5.1e-35	1.7e-30	1e-38	1.7e-27
END		141	140	140	120	152	134
START AA		4		4	2	6	17
CHAIN		Y	A	A	D	Q	
PDB ID		1bu9	lihb	lihb	likn	lika	Ішуо
SEQ ID NO:		397	397	397	397	397	397

	<u>.</u>	NO NO	NO NO		PCI		SOE/D1E	
PDB annotation	ACETYLATION, NMR, ANK- REPEAT	COMPLEX (TRANSCRIPTION REG/ANK REPEAT) COMPLEX (TRANSCRIPTION REGULATION/ANK REPEAT), ANKYRIN 2 REPEAT, HEI IX	COMPLEX (TRANSCRIPTION REG/ANK REPEAT) COMPLEX (TRANSCRIPTION REG/ULATION/ANK REPEAT), ANKYRIN 2 REPEAT HELIX			METAL TRANSPORT CALMODULIN, HIGH RESOLUTION, DISORDER	SERINE/THREONINE FOR PROTEIN KINASE TRANSFERASE, SERINE/THREONINE ONCOGENE, ZINC, ATP-BINDING, PHORBOL-ESTER BINDING	SERINE/THREONINE PROTEIN KINASE
Compound		NF-KAPPA-B P65; CHAIN: A, C; NF-KAPPA-B P50; CHAIN: B, D; I-KAPPA-B- ALPHA; CHAIN: B, F;	NF-KAPPA-B P65; CHAIN: A, C; NF-KAPPA-B P50; CHAIN: B, D; I-KAPPA-B- ALPHA; CHAIN: E, F;	CALCTUM-BINDING PROTEIN CALMODULIN COMPLEXED WITH CALMODULIN-BINDING DOMAIN OF ICDM 3 CALMODULIN- DEPENDENT PROTEIN KINASE II ICDM 4	CALCIUM-BINDING PROTEIN CALMODULIN (VERTEBRATE) ICLL 3	CALMODULIN; CHAIN: A;	RAF-1; CHAIN: NULL;	RAF-1; CHAIN: NULL;
SEQ FOLD score								·
PMF score		1.00	1.00	0.37	0.35	0.15	0.58	0.47
Verify score	٠	0.59	0.51	-0.27	-0.42	-0.42	0.33	0.19
Psi Blast		1.7e-30	1e-38	6.8e-33	1.2e-36	6.8e-35	1.2e-06	2.6e-15
END		120	152	239	239	239	310	312
START AA		2	6	91	16	68	261	261
CHAIN		时	E	¥		А		
PDB ID		lnfi	1nfi	1cdm	Icli	lexr	lfag	lfaq
SEQ ID NO:		397	397	400	400	400	400	400

Table

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PDB annotation	TRANSFERASE, SERINE/THREONINE- PROTEIN KINASE, 2 PROTO- ONCOGENE, ZINC, ATP- BINDING, PHORBOL-ESTER BINDING	SERINETHREONINE PROTEIN KINASE TRANSFERASE, SERINETHREONINE- PROTEIN KINASE, 2 PROTO- ONCOGENE, ZINC, ATP- BINDING, PHORBOL-ESTER BINDING	SERINE/THREONINE PROTEIN KINASE TRANSFERASE, SERINE/THREONINE- PROTEIN KINASE, 2 PROTO- ONCOGENE, ZINC, ATP- BINDING, PHORBOL-ESTER BINDING	PHOSPHOTRANSFERASE	PHOSPHOTRANSFERASE (PHOSPHOTRANSFERASE	CALCIUM-BINDING PROTEIN RAT BRAIN PKC- () G; CALCIUM-BINDING PROTEIN, PROTEIN KINASER C, PKC, TRANSFERASE	CALCTUM-BINDING PROTBIN RAT BRAIN PKC- C G; CALCTUM-BINDING PROTBIN, PROTBIN KINASER C, PKC, TRANSFERASE	CALCIUM-BINDING
Compound		RAF-1; CHAIN: NULL;	RAF-1; CHAIN: NULL;	PROTEIN KINASE C DELTA TYPE; 1PTQ 4	PROTEIN KINASE C DELTA TYPE; 1PTQ 4	PROTEIN KINASE C DELTA TYPE; 1PTQ 4	PROTEIN KINASE C, GAMMA TYPE, CHAIN: NULL;	PROTEIN KINASE C, GAMMA TYPE; CHAIN: NULL;	TROPONIN C; 1TNX 4
SEQ FOLD score									
PMF		0.01	0.19	0.19	0.19	0.31	0.25	0.25	0.11
Verify		-0.42	-0.37	-0.16	-0.16	0.17	-0.36	-0.22	-0.56
Psi Blast		2e-07	6.8e-06	6.6e-17	6.8e-10	3.4e-17	1e-09	1.7e-18	1.7e-25
END AA		374	374	310	310	374	310	316	239
START AA		334	336	261	761	325	261	261	91
CHAIN									
PDB ID		lfaq	lfaq	· 1ptq	lptq	1ptq	1tbn	1tbn	1tmX
SEQ ID NO:		400	400	400	400	400	400	400	400

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PDB annotation	PROTEIN EF-HAND 1TNX 14		CALMODULIN, CALCIUM BINDING, HELIX-LOOP- HELIX, SIGNALLING, 2 COMPLEX(CALCIUM- BINDING PROTEIN/PEPTIDE)		OXIDOREBUCTASB PDZ DOMAIN, NNOS, NITRIC OXIDE SYNTHASB	Corne do Amortica do Day	DOMAIN, NNOS, NITRIC	OXIDE SYNTHASE	OXIDOREDUCTASE PDZ	DOMAIN, NNOS, NITRIC	OXIDE SYNTHASE	PEPTIDE RECOGNITION	PEPTIDE RECOGNITION, PROTEIN LOCALIZATION	PEPTIDE RECOGNITION	PROTEIN LOCALIZATION	PEPTIDE RECOGNITION	PROTEIN LOCALIZATION	KINASE HCASK, GLGF	REPEAT, DHR; PDZ	SYNDECAN, RECEPTOR	CLUSTERING, KINASE	KINASE HCASK, GLGF REPEAT, DHR; PDZ DOMAIN, NEUREXIN,
Compound	CHAIN: NULL; 1TNX 5	CONTRACTILE SYSTEM PROTEIN TROPONIN C 1TOP 3	CALMODULIN; CHAIN: A; RS20; CHAIN: B;		NEURONAL NITRIC OXIDE SYNTHASE; CHAIN: A; HEPTAPEPTIDE; CHAIN: B;	TOTAL ATTENDED	NEURONAL MILKIC OXIDE SYNTHASE;	CHAIN: A; HEPTAPEPTIDE: CHAIN: B:	NEURONAL NITRIC	OXIDE SYNTHASE;	CHAIN: A; HEPTAPEPTIDE; CHAIN: B;	PSD-95; CHAIN: A; CRIPT;	CHAIN: B;	PSD-95; CHAIN: A; CRIPT;	CHAIN: B;	PSD-95; CHAIN: A; CRIPT;	Chally: B;	HCASK/LIN-2 PROTEIN;	CHAIN: A, B;			HCASK/LIN-2 PROTEIN; CHAIN: A, B;
SEQ FOLD score											-								•			
PMF		0.06	0.22	,	1:00	,,,,	0.00		0.99			0.99		1.00		0.39		0.92				0.39
Verify		0.07	-0.41		0.64	9.0	0.10		0.29			0.26		0.61		0.14		0.51				-0.16
Psi Blast		3.4e-27	6.8e-35		3.3e-15	,,	9.9e-10		9.9e-17			5.1e-07		1.7e-16		1.4e-17		3.3e-15				3.3e-14
END		239	240		. 290	200	893		157			288		166		8		293				893
START		91	88		222	į	818		84			232		75		812		222				820
CHAIN			V		∢		∢		4			A		Ą		Ą		Ą				Ą
PDB		Itop	lvrk		1b8q		1589		1b8q	,		1bes		1be9		1be9		1kwa				1kwa
SEQ ID		400	400		403		403		403			403		403		403		403				403

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PDB annotation	SYNDECAN, RECEPTOR CLUSTERING, KINASE	KINASE HCASK, GLGF REPEAT, DHR; PDZ DOMAIN, NEUREXIN, SYNDECAN, RECEPTOR	SIGNAL TRANSDUCTION HDLG, DHR3 DOMAIN; SIGNAL TRANSDUCTION, SH3 DOMAIN, REPEAT	SIGNAL TRANSDUCTION HDLG, DHR3 DOMAIN; SIGNAL TRANSDUCTION, SH3 DOMAIN, REPEAT	SIGNAL TRANSDUCTION HDLG, DHR3 DOMAIN; SIGNAL TRANSDUCTION, SH3 DOMAIN, REPEAT	OXIDOREDUCTASE BETA- FINGER	OXIDOREDUCTASE BETA- TO FINGER	MEMBRANE PROTEIN/OXIDOREDUCTAS E BETA-FINGER, HETERODIMER	MEMBRANE PROTEIN/OXIDOREDUCTASE B BETA-FINGER, HETERODIMER
Compound		HCASKLIN-2 PROTEIN; CHAIN: A, B;	HUMAN DISCS LARGE PROTEIN; CHAIN: NULL;	HUMAN DISCS LARGE PROTEIN; CHAIN: NULL;	HUMAN DISCS LARGE PROTEIN; CHAIN: NULL;	NEURONAL NITRIC OXIDE SYNTHASE (RESIDUES 1-130); CHAIN: A;	NEURONAL NITRIC OXIDE SYNTHASE (RESIDUES 1-130); CHAIN: A;	ALPHA-1 SYNTROPHIN (RESIDUES 77-171); CHAIN: A; NEURONAL NITRIC OXIDE SYNTHASE (RESIDUES 1-130); CHAIN: B;	ALPHA-1 SYNTROPHIN (RESIDUES 77-171); CHAIN: A; NEURONAL NITRIC OXIDE SYNTHASE (RESIDUES 1-130); CHAIN: B;
SEQ FOLD score									
PMF score		0.94	1.00	0.34	0.98	0.87	0.93	1.00	0.35
Verify score		0.41	0.16	0.05	0.66	0.38	0.67	0.91	-0.18
Psi Blast		1.3e-18	9.9e-12	1.7e-16	1.7e-15	6.6e-15	2e-18	6.6e-17	1.7e-16
END AA		168	273	904	166	303	170	289	905
START AA			222	818	833	222	84	222	815
CHAIN		Ą				4	Ą	∢	¥
PDB ID		1kwa	1pdr	lpdr	1pdr	1qau	Iqau	lqav	Iqav
SEQ ID NO:		403	403	403	403	403	403	403	403

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PDB annotation	MEMBRANE PROTEIN/OXIDOREDUCTAS E BETA-FINGER, HETERODIMER	MEMBRANE PROTEIN/OXIDOREDUCTAS B BETA-FINGER, HETERODIMER	MEMBRANE PROTEIN/OXIDOREDUCTAS B BETA-FINGER, HETERODIMER	PEPTIDE RECOGNITION PSD-95; PDZ DOMAIN, NEURONAL NITRIC OXIDE SYNTHASE, NMDA RECEPTOR 2 BINDING	PEPTIDE RECOGNITION PSD-95; PDZ DOMAIN, NEURONAL NITRIC OXIDE SYNTHASE, NMDA **RECEPTOR 2 BINDING	PEPTIDE RECOGNITION PSD-95; PDZ DOMAIN, NEURONAL NITRIC OXIDE SYNTHASE, NMDA RECEPTOR 2 BINDING	PEPTIDE RECOGNITION PSD-95; PDZ DOMAIN, NEURONAL NITRIC OXIDE SYNTHASE, NMDA RECEPTOR 2 BINDING	HYDROLASE PDZ DOMAIN, 🖺
Compound	ALPHA-1 SYNTROPHIN (RESIDUES 77-171); CHAIN: A; NEURONAL NITRIC OXIDE SYNTHASE (RESIDUES 1-130); CHAIN: B;	ALPHA-1 SYNTROPHIN (RESIDUES 77-171); CHAIN: A; NEURONAL NITRIC OXIDE SYNTHASE (RESIDUES 1-130); CHAIN: B:	ALPHA-1 SYNTROPHIN (RESIDUES 77-171); CHAIN: A; NEURONAL NITRIC OXIDE SYNTHASE (RESIDUES 1-130); CHAIN: B:	POSTSYNAPTIC DENSITY PROTEIN 95; CHAIN: A;	POSTSYNAPTIC DENSITY PROTEIN 95; CHAIN: A;	POSTSYNAPTIC DENSITY PROTEIN 95; CHAIN: A;	POSTSYNAPTIC DENSITY PROTEIN 95; CHAIN: A;	TYROSINE PHOSPHATASE
SEQ FOLD score								
PMF	1.00	1.00	0.99	0.1	1.00	0.99	1.00	1.00
Verify score	0.53	0.64	0.77	0.77	0.22	0.07	0.72	0.45
Psi Blast	6.6e-15	9.9e-20	5.1e-18	2.3e-14	2.3e-16	1.2e-14	1.3e-17	1.6e-14
END	893	165	168	289	893	106	165	289
START AA	. 918	81	81	222	816	819	81	222
CHAIN	4	¥	∀	∢	4	A	Ą	Ą
PDB ID	lqav	Iqav	lqav	1qic	1qic	1qlc	1qlc	3pdz
SEQ ID NO:	403	403	403	403	403	403	403	403

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PDB annotation	HUMAN PHOSPHATASE, HPTP1E, PTP-BAS, SPECIFICITY 2 OF BINDING	HYDROLASE PDZ DOMAIN, HUMAN PHOSPHATASE,	HPIPIE, PIP-BAS, SPECIFICITY 2 OF BINDING	HYDROLASE PDZ DOMAIN, HUMAN PHOSPHATASE,	HPTP1E, PTP-BAS, SPECIFICITY 2 OF BINDING	HYDROLASE PDZ DOMAIN, HUMAN PHOSPHATASE,	HPTP1E, PTP-BAS, SPECIFICITY 2 OF BINDING	HYDROLASE PDZ DOMAIN,	HPTPIE, PTP-BAS,	SPECIFICITY 2 OF BINDING		COMPLEX (ZINC FINGER/DNA) COMPLEX	-	PROTEIN	COMPLEX (ZINC #	(ZINC FINGER/DNA), ZINC (FINGER, DNA-BINDING	PROTEIN	COMPLEX (ZINC	(ZINC FINGER/DNA), ZINC	FINGER, DNA-BINDING	PROTEIN	COMPLEX (ZINC
Compound	(PTP-BAS, TYPB 1); CHAIN: A;	TYROSINE PHOSPHATASE (PTP-BAS, TYPE 1); CHAIN:	Α;	TYROSINE PHOSPHATASE (PTP-BAS, TYPE 1); CHAIN:	À;	TYROSINE PHOSPHATASE (PTP-BAS, TYPE 1); CHAIN:	A ;	TYROSINE PHOSPHATASE	A;			QGSR ZINC FINGER PEPTIDE; CHAIN: A;	OLIGONUCLEOTIDE	BINDING STIE; CHAIN: B, C;	QGSR ZINC FINGER	PEPTIDE; CHAIN: A; DUPLEX	OLIGONUCLEOTIDE	BINDING SITE; CHAIN: B, C;	OGSR ZINC FINGER	DUPLEX	OLIGONUCLEOTIDE	BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER
SEQ FOLD score																							
PMF score		0.88		0.41		9.76		1.00				0.68			1.00				1.00				1.00
Verify score		0.02		0.26		0.37		0.27				-0.55			0.33		_		-0.04			,	0.49
Psi Blast		8.5e-09		1.7e-16		1.7e-15		1.3e-16				1.7e-23			3.4e-29				3.3e-33				3.4e-31
END AA		296		206		171		170				222			250				251			•	306
START AA		233		814	•	81	·	84		-		147			0/1				171				226
CHAIN		V		Ą		¥		Ą				∢			¥				Ą				Ą
PDB ID		3pdz		3pdg		3pqz		zpdg				1alh			lalh				lalh				lalh
SEQ ID NO:		403		403	•	403		403				2 04			404				404				404

Table

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PDB annotation	FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER) FINGER DIA PROTEIN-DNA PROTEIN INTERACTION, PROTEIN FIN
Compound	PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCI EOTIDE BINDING SITE; CHAIN: B, C;	DNA; CHAIN: A, B, D, B; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;
SEQ FOLD score			80.11		·		
PMF		1.00		1.00	1.00	0.11	0.03
Verify score		0.43		0.12	0.38	-0.09	-0.26
Psi Blast		9.96-34	9.96-34	3.3e-32	9.96-31	3.46-19	6.8e-36
AA END		307	308	334	362	393	194
START AA		226	226	254	282	310	118
CHAIN		∢	∢	¥	¥	¥	ပ
PDB ID		lalh	laih	lalh	laIh	laih	lmey
SEQ ID NO:		404	404	404	404	404	404

Table 5

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PDB annotation	DESIGN, 2 CRYSTAL STRUCTURB, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROFEREN PANA	INTERACTION, PROTEIN	DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER	PROTEIN-DNA	INTERACTION, PROTEIN	DESIGN, 2 CRYSTAL	SIRUCIONE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC	FINGER/DNA) ZINC FINGER,	PROTEIN-DINA	DESIGN 2 COVETAI	STRIICTIBE COMPLEX	(ZINC FINGER/DNA)		FINGER/DNA) ZINC FINGER,	PROTBIN-DNA	INTERACTION, PROTEIN	DESIGN, 2 CRYSTAL	STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC	PROTEIN-DNA	INTERACTION, PROTEIN	DESIGN, 2 CRYSTAL	STRUCTURE, COMPLEX	(A)	COMPLEX (ZINC
Compound		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER	ANOTHER, CIMEN. C. F. G.			DNA; CHAIN: A, B, D, E;	PROTEIN; CHAIN: C, F, G;				DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;				DNA: CHAIN: A B D E.	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;					DNA; CHAIN: A, B, D, E;	PROTEIN: CHAIN: C. F. G.				4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4	DNA; CHAIN: A, B, D, E;
SEQ FOLD score																		·							•				,000	98.20
PMF		0.86				1.00					1.00						8	8						1.00						
Verify		-0.06				0.08				·	0.35						0 38	2						0.32						
Psi Blast		1.76-42				1.7e-50					3.4e-51						680.57	7000						5.1e-47					,	3.164/
END		222				250					278						306	3						362						200
START AA	·	146				169					197						225	3						253						281
CHAIN		၁				ບ					ည						ر)						 ပ					,	5
PDB ID		Ітеу				lmey					lmey			-) Tabe	- Carrier I						Imey						Imey
SEQ ID NO:		404				404					404						404	·						404						404

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PDB annotation	FINGER,DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER,DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER,DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) TFIIIA; 5S GENE; NMR, TFIIIA, PROTEIN, DNA, TRANSCRIPTION FACTOR, 5S RNA 2 GENE, DNA BINDING PROTEIN, ZINC FINGER, COMPLEX 3 (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION) REGULATIONDNA) COMPLEX (TRANSCRIPTION) REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INTIATION, ZINC FINGER PROTEIN
Compound	CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	TRANSCRIPTION FACTOR IIIA; CHAIN: A; SS RNA GBNB; CHAIN: B, P;	TFIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	TFIIIA; CHAIN: A, D; SS RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;
SEQ FOLD score			60.10	114.59	
PMF score		0.04			1.00
Verify		-0.12			-0.07
Psi Blast		8.5e-38	1.7e-18	3.46-38	3.4e-38
END AA		393	300	337	315
START		309	225	167	170
CHAIN		U U	4	V	4
PDB EDB		lmey	1453	1tf6	1tf6
SEQ ID NO:		404	404	404	404

PDB annotation	COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA	POLYMERASE III, 2	INTITATION, ZINC FINGER	PROTEIN	COMPLEX (TRANSCRIPTION	YANG 1; TRANSCRIPTION	INITIATION, INITIATOR	ELEMENT, YY1, ZINC2	PROTEIN PROCESSIFION 2	COMPLEX (TRANSCRIPTION	REGUL ATION/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA) YING-	YANG 1; TRANSCRIPTION	INITIATION, INITIATION	ELEMENT, YY1, ZINC 2	FINGER PROTEIN, DNA-	PROTEIN RECOGNITION, 3	COMPLEX (TRANSCRIPTION	REGULATION/DNA)	COMPLEX (TRANSCRIPTION)	VANG 1: TP ANSCRIPTION	INITIATION, INITIATOR	ELEMENT, YY1, ZINC 2	FINGER PROTEIN, DNA-	PROTEIN RECOGNITION, 3	COMPLEX (TRANSCRIPTION	REGULATION/DNA)	COMPLEX (TRANSCRIPTION PROFILE)	YANG I; TRANSCRIPTION	INITIATION, INITIATOR FI
Compound	TFIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE;	CHAIN: B, C, E, F;				YY1; CHAIN: C; ADENO-	ASSOCIATED VIKUS FS INITIATOR ELEMENT	DNA; CHAIN: A, B;					YY1; CHAIN: C; ADENO-	ASSOCIATED VIRUS P5	INITIATOR ELEMENT	DNA; CHAIN: A, B;					TANI CITABLE AND INC.	A SEOCIATION AMERICAN	INTERIOR HI PARINT	DNA; CHAIN: A. B:						YYI; CHAIN: C; ADENO- ASSOCIATED VIRIES P5	INITIATOR ELEMENT	DNA; CHAIN: A, B;
SEQ FOLD score																																
PMF	0.62					0.98							1.00					_			00.	7.0	•							1.00		
Verify	-0.04		•			0.03							0.05								21.0	07.0								0.23		
Psi Blast	5.1e-34					16-31							1.3e-39								1 20 34	1.25.7					•			3.3e-43		
AA AA	403					250							278								270	0/1								335		
START AA	254					148			-			•	174						,		177									223		1
CHAIN	V			·		ບ		•					ن					٠.	_		ļ	 ,										
PDB ID	1tf6			-		lubd							1ubd							•	1hd	3								pani		
SEQ ID NO:	404					4 4							\$							•	707	<u> </u>						-		\$		

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				limate Named" walless "to grand range Handle to		p. Harp, ram
PDB annotation	ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA- PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATIONDNA) YING- YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA- PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATIONDNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 (COMPLEX (TRANSCRIPTION) REGULATION/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-FINGER GIL; GIL; ZINC FINGER, COMPLEX (DNA-BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-
Compound		YYI; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	ZINC FINGHR PROTRIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA;
SEQ FOLD score		87.63		·		
PIMF score			0.89	1.00	0.41	1.00
Verify score			0.04	0.0- 80.0-	-0.06	0.43
Psi Blast		3.36-43	5.1e-32	3.3e-38	1.2e-31	3.4e-33
END			362	362	249	305
START	,	225	233	251	126	177
CHAIN ID		ပ	ນ	၁	¥	A
PDB ID			Jubd	pqnI	2gli 7	2gli /
SEQ ID NO:		404	404	404		404

			7	<u> </u>	PL		502/01	223
PDB annotation	FINGER, COMPLEX (DNA-BINDING)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE- FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE- FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	POTASSIUM CHANNELS POTASSIUM CHANNELS, TETRAMERIZATION DOMAIN, X-RAY 2 STRUCTURE, APLYSIA	SIGNALING PROTEIN VOLTAGE-GATED POTASSIUM CHANNEL, ASSEMBLY DOMAIN, TETRAMER	METAL TRANSPORT ION ' CHANNEL, OXIDOREDUCTASE, BETA SUBUNIT	SIGNALING PROTEIN VOLTAGE-GATED POTASSIUM CHANNEL, TETRAMERIZATION DOMAIN, 2 INTRACELLULAR GATE, TETRAMER	PROTON TRANSPORT POTASSIUM CHANNELS, TETRAMERIZATION
Compound	CHAIN: C, D;	ZINC FINGER PROTEIN GLI1; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	POTASSIUM CHANNEL KV1.1; CHAIN: NULL;	KV1.2 VOLTAGE-GATED POTASSIUM CHANNEL; CHAIN: A, B, C, D, B, F, G, H;	KV BETA2 PROTEIN; CHAIN: A; POTASSIUM CHANNEL KV1.1; CHAIN: E;	KV1.2 VOLTAGE-GATED POTASSIUM CHANNEL; CHAIN: A, B, C, D;	POTASSIUM CHANNEL KV1.1; CHAIN: A;
SEQ FOLD score		98.48						•
PMF score			1.00	0.76	0.92	0.98	0.98	0.98
Verify score			0.21	0.31	0.22	0.27	0.34	0.16
Psi Blast		3.4e-33	6.8e-30	1.4e-27	3.46-26	3.4e-27	3.4e-29	1.7e-29
END			361	135	135	136	149	150
START AA	·	197	205	51	51	50	51	51
CHAIN		Ą	¥	į:	¥	B	¥	Ą
PDB ID		2gli	2gli	1a68	ldsx	lexb	vbpl.	ltld
SEQ ID NO:		404	404	405	405	405	405	405

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PDB annotation	DOMAIN, X-RAY STRUCTURE, 2 APLYSIA KV1.1, PROTON TRANSPORT	POTASSIUM CHANNEL POTASSIUM CHANNEL, TETRAMERIZATION DOMAIN, MOLECULAR 2 RECOGNITION, ZINC- BINDING		T-CELL SURFACE GLYCOPROTEIN IMMUNOGLOBULIN FOLD, TRANSMEMBRANE, GLYCOPROTEIN, T-CELL, 2	MHC, LIPOPROTEIN, T-CELL SURFACE GLYCOPROTEIN	GROWTH FACTOR/GROWTH FACTOR RECEPTOR FGF.	FGFR, IMMUNOGLOBULIN- LIKE, SIGNAL	TRANSDUCTION, 2 DIMERIZATION, GROWTH	FACTOR/GROWITH PACTOR T	GROWTH FACTOR/GROWTH FACTOR RECEPTOR FGF, "	FGFR, IMMENOGLOBULIN-	TRANSDUCTION, 2	DIMERIZATION, GROWTH	RECEPTOR	VIRUS/VIRAL PROTBIN, RECEPTOR CD155, PVR, FUMAN POLIOVIRUS,	ELECTRON MICROSCOPY, 2 POLIOVIRUS-RECEPTOR	COMPLEX, VIRUS/VIRAL
Compound		POTASSIUM CHANNEL PROTEIN SHAW; CHAIN: NULL;		T-CELL SURFACE GLYCOPROTEIN CD4; CHAIN: NULL;		FIBROBLAST GROWTH FACTOR 2; CHAIN: A, B;	FIBROBLAST GROWTH FACTOR RECEPTOR 1:	CHAIN: C, D;		FIBROBLAST GROWTH FACTOR 2; CHAIN: A, B;	FIBROBLAST GROWTH	CHAIN: C, D;			POLIOVIRUS RECEPTOR; CHAIN: R; VP1; CHAIN: 1; VP2; CHAIN: 2; VP3;	CHAIN: 3; VP4; CHAIN: 4;	
SEQ FOLD score													-				
PMF score		0.52		7.0		950				0.30					1.00		
Verify score		0.55		0.36		0.33				0.37					0.26		
Psi Blast		1.7e-33		9.9e-12		1.3e-08				2e-08					2e-08		
END		151		138		145				145					136		
START AA	·	. 05		20		50			:	- 20					47		
CHAIN						ပ				Ω .		·			~		
PDB ID		3kvt		lody		Icvs				lcvs					Idgi		
SEQ ID NO:		405		407		407				407					407		

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PDB annotation	PROTEIN, RECEPTOR					BELONGING TO THE I-SET 2	SUBGROUP WITHIN IG-LIKE	A CONTRACTILE PROTEIN	IMMUNOGLOBULIN FOLD,	BETA BARREL		CONNECTIN, NEXTINS;	Car ACOBOTTEIN	TRANSMEMBRANE	REPEAT BRAIN 2	IMMUNOGLOBULIN FOLD.	ALTERNATIVE SPLICING,	SIGNAL, 3 MUSCLE	PROTEIN	TITIN	•	M3	S. III	'URB)	1,1	GLYCOPROTEIN CD4;		O TOO TO TO TO TO TO TO TO TO TO TO TO T	GLICOFROIEIN, 1-CELL, 2 (1)	POLYMORPHISM	TEIN 40; CHAPERONE BETA SHEETS.		TEIN 40; CHAPERONE BETA SHEETS 4	
Compound		FIBROBLAST GROWTH	FACTOR 1; CHAIN: A, B;	FIBROBLASI GROWIH	FACTOR RECEPTOR 1;	CHAIN: C, D;		TELOKIN; CHAIN: A			TITIN; CHAIN: NULL;									MUSCLE PROTEIN TITIN	CIVI STROTONI	(CONNECTIN) 1TNM 3	(NMK, MINIMIZED	AVERAGE STRUCTURE)	ITINM 4 ITINM 58	T-CELL SURFACE	CHAIN A B.	CITATIV: A, D,	-		HEAT SHOCK PROTEIN 40;	CHAIN: A;	HEAT SHOCK PROTEIN 40; CHAIN: A:	HUMAN HSP40; CHAIN:
SEQ FOLD score		٠					-						•		÷																			
PMF score		0.57		_				1.00		,	66'0									96'0						0.46					1.0		00'1	1.00
Verify score		0.46						0.19			0.53				-					0.92						-0.07					0.34		0.26	0.59
Psi Blast		26-08						6.6e-08			6.8e-09							-		6.8e-09						6.ee-09					3.3e-53		6.8e-33	2.3e-29
END		136						138			136									921						138					310		310	75
START AA		47			_			47			43									43						49					140		145	1
CHAIN		၁						≺																		∢					¥		Ą	
PDB TD		levt						1fhg			lnct									Itmm						lwio					163g	,	1c3g	Ihdj
SEQ ID NO:		407						407			407									407						407					408		408	408

rable:

PDB annotation	HDJ-1; MOLECULAR CHAPERONE	MOLECULAR CHAPERONE HDJ-1; MOLECULAR CHAPERONE	MOLECULAR CHAPERONE HDJ-1; MOLECULAR CHAPERONE	2 1 2 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	CONTRACTILE LIM DOMAIN, CRP, NMR,	DIFFERENTIATION, CONTRACTILE	COMPLEX (ZINC FINGER/DNA) ZINC FINGER,	PROTEIN-DNA	INTERACTION, PROTEIN	STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER,	PROTEIN-DNA	INTERACTION, PROTEIN	STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER U	PROTEIN-DNA	DESIGN 2 CRYSTAI	STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC FINGER T	PROTEIN-DNA NTERACTION, PROTEIN N
Compound	NULL;	HUMAN HSP40; CHAIN: NULL;	HUMAN HSP40; CHAIN: NULL;	A. A. C. C. C. C. C. C. C. C. C. C. C. C. C.	CRP1; CHAIN: A;		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;	-			DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;				DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G,				DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;
SEQ FOLD score		93.05		3, 3,	03.60																		
PMF score	·		1.00				0.96					1.00					1.00					1.00	
Verify score			0.75				0.19					0.34					0.78					0.10	
Psi Blast		2.3e-29	3.4e-25		6.0e-15		1e-45					3.4e-46					1.2e-47					5.1e-48	
END AA		75	76	9	409		191			-		219		_			247					275	
START AA		1	1	, 50	4/7		110					138					166					194	
CHAIN	·				∢		ي					ပ					ပ				·	ບ	
PDB TO		1hdj	1hdj	1011	1081		Ітеу					1теу					1mey					1теу	
SEQ ID NO:		408	408	9.5	410		410					410					410					410	

Fable :

PDB annotation	DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN, DNA	INTERACTION, PROTEIN	DESIGN, 2 CRYSTAL STRICTURE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC	FINGER/DNA) ZINC FINGER; PROTEIN-DNA	INTERACTION, PROTEIN	DESIGN, 2 CRYSTAL	STRUCTURE, COMPLEX	(ZINC FINGER/DIVA)	FINGER/DNA) ZINC FINGER,	PROTEIN-DNA	INTERACTION, PROTEIN	DESIGN, 2 CRYSTAL	STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC	FINGER/DNA) ZINC FINGER,	PROTEIN-DNA	INTERACTION, PROTEIN	DESIGN, 2 CRYSTAL	STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC	FINGER/DNA) ZINC FINGER;	PROTEIN-DNA	INTERACTION, PROTEIN	 ZINCTIONE, COMPLEA 18	COMPLEX (ZINC
Compound		DNA; CHAIN: A, B, D, B; CONSENSUS ZINC FINGER PROTEIN: CHAIN: C B G:				DNA; CHAIN: A, B, D, B;	CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;				T C C T T T T T T T T T T T T T T T T T	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, P, G;					DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;	,				DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;			DNA; CHAIN: A, B, D, E;
SEQ FOLD score																														95.94
PMF score		1.00	•			1.00						00.1						1.00							00'1					
Verify		90.0				-0.19					2,0	cr.o		,				0.13							0.41					
Psi Blast		1.7e-49				1.7e-49						1./6-50						1.7e-50							6.8e-51					6.8e-51
END		303				331			•		0,00	959						387							415					416
START		222				250					010	8/7						306						i	334					334
CHAIN		၁				၁					,	ر-						ပ							၁				-	C
PDB ID		1mey	_			1mey						Iney						1mey							1mey					Imey
SEQ ID NO:		410				410					Š	410						410							410					410

Table

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PDB annotation	FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX ZTNC HINGER DNA)	COMPLEX (ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX CTNC FINGER, DNA PONTEIN DIGGRESS (COMPLEX CTNC FINGER, DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX CZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER(F) PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX	COMPLEX (ZINC EPINGER FINGER) PROTEIN-DNA INTERACTION, PROTEIN FINDERICAL DESIGN, 2 CRYSTAL FINDERICAL FINDERI
Compound	CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, B; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, B; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;
SEQ FOLD score						
PMF		1.00	1.00	1.00	0.13	0.05
Verify		0.26	0.51	0.26	-0.32	-0.41
Psi Blast		1.2e-50	3.4e-50	1.5e-33	6.8e-43	1.7e-44
END AA		443	471	475	135	163
START AA		362	390	418	54	82
CHAIN ID		U	U	O	၁	ນ
PDB ID		Imey	lmey	1 mey	1mey	1mey
SEQ ID NO:	·	410	410	410	410	410

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PDB annotation	STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INTIATION, ZINC FINGER	COMPLEX (TRANSCRIPTION REGULATIONDNA) COMPLEX (TRANSCRIPTION REGULATIONDNA), RNA POLYMERASE III, 2 TRANSCRIPTION INTIATION, ZINC FINGER PROTTEN	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASB III, 2 TRANSCRIPTION INITIATION, ZINC FINGER	COMPLEX (TRANSCRIPTION REGULATION/DNA) COMPLEX (TRANSCRIPTION REGULATION/DNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATIONDNA) COMPLEX (TRANSCRIPTION REGULATIONDNA), RNA [1] POLYMERASE III, 2
Compound		TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, B, F;	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, B, F;	TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, B, F;
SEQ FOLD score				104.78		
PMF		0.66	0.83		1.00	0.43
Verify score		0.08	-0.09		0.01	-0.46
Psi Blast		5.1e-36	1e-38	1,7e-36	1.7e-36	3.46-34
END AA		256	368	474	473	205
START AA		111	223	306	335	55
CHAIN		∀	∢	∢	∢	∢
PDB ID		1446	1476	1tf6	11f6	1466
SEQ ID NO:		410	410	410	410	410

 Table

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PDB annotation	TRANSCRIPTION INITIATION, ZINC PINGER PROTEIN	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING- YANG 1; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA- PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION TO INITIATION, INITIATOR FLEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-FROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION) REGULATION/DNA) YING- [1] YANG 1; TRANSCRIPTION , INITIATION, INITIATOR , ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA- H PROTEIN RECOGNITION, 3 [1] COMPLEX (TRANSCRIPTION] REGULATION/DNA)
Compound		YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INTIATOR ELEMENT DNA; CHAIN: A, B;	YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;	YY1; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO-ASSOCIATED VIRUS PS INITIATOR ELEMENT DNA; CHAIN: A, B;
SEQ FOLD score	·		88.64		
PMF score		1.00		1.00	1.00
Verify score		0.20		0.02	60:0
Psi Blast		2e-38	3.3e-50	3.3e-50	6.6e-56
END AA			248	248	304
START		120	139	141	193
CHAIN		v	υ	်	ပ
PDB DD		1ubd	pqm1	lubd	1ubd
SEQ ID NO:		410	410	410	410

Table 5

PDB annotation	COMPLEX (TRANSCRIPTION REGULATIONDNA) YING- YANG I; TRANSCRIPTION INITIATION, INITIATOR ELEMENT, YY1, ZINC 2	FINGER PROTEIN, DNA- PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING- XANG 1. TE ANSCRIPTION	INTIATION, INTIATOR ELEMENT, YYI, ZINC 2	FINGER PROTEIN, DNA-	COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION	YANG 1; TRANSCRIPTION	INITIATION, INITIATOR FILEMENT YYL ZINC 2	FINGER PROTEIN, DNA-	PROTEIN RECOGNITION, 3 [7]	COMPLEX (TRANSCRIPTION) REGULATION/DNA)	COMPLEX (TRANSCRIPTION	YANG 1; TRANSCRIPTION ()	HITTATION, INITIATOR	FINGER PROTEIN, DNA-	PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION	REGULATION/DNA)	7 .	TANG I; INCHASCRICTION
Compound	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;		YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS PS	DNA; CHAIN: A, B;			YY1; CHAIN: C; ADENO-	INITIATOR ELEMENT	DNA; CHAIN: A, B;			•	YY1; CHAIN: C; ADENO-	ASSOCIATED VIRUS PS INITIATOR ELEMENT	DNA; CHAIN: A, B;				YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5	INITIATOR ELEMENT
SEQ FOLD score																				
PMF score	1.00		1.00				1.00						0.70						1.00	
Verify score	-0.02		-0.12				-0.07				•		0.02						-0.11	
Psi Blast	1e-34	·	1.2e-35			-	3.3e-53						2e-53						1.2e-35	
END	303		331				359						415		•				387	
START AA	202		224				249		•				276						286	
CHAIN	ပ		ວ				၁	 · -					၁						O	
PDB ID	lubd		1ubd				Iubd						1ubd						lubd	1
SEQ ID NO:	410		410				410						410						410	

PDB annotation	INITIATION, INITIATOR ELEMENT, YYI, ZINC 2 FINGER PROTEIN, DNA- PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATOR HIMOER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING- YANG 1; TRANSCRIPTION HINTIATION, INITIATOR ELEMENT, YY1, ZINC2 FINGER PROTEIN, DNA-
Compound	DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INTIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;
SEQ FOLD score					
PMF score		1.00	1.00	1.00	1.00
Verify		0.12	0.29	0.29	0.20
Psi Blast		3.3e-54	8.5e-36	2e-51	16-34
END AA		443	443	470	471
START AA		332	339	360	370
CHAIN		U	U	ပ	υ
POB		lubd	lubd	1ubd	lubd
SEQ ID		410	410	410	410

Table !

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PDB annotation	PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING- YANG 1; TRANSCRIPTION INITIATION INITIATOR	ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA- PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YÏNG- YANG 1; TRANSCRIPTIQN INITIATION, INITIATOR	ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-	COMPLEX (TRANSCRIPTION BEGIN ATTOMONA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE.	FINGER, COMPLEX (DNA-	BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE-	FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA-	BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING 19 PROTFINIONA) RIVE.	FINGER GLI, GLI, ZINC	FINGER, COMPLEX (DNA-	COMPLEX (DNA-BINDING IU	PROTEIN/DNA) FIVE-
Compound		YYI; CHAIN: C; ADBNO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA: CHAIN: A. B:		YY1; CHAIN: C; ADENO-ASSOCIATED VIRUS PS INITIATOR ELEMENT ODNA; CHAIN: A, B;			ZINC FINGER PROTEIN GLII; CHAIN: A; DNA;	CHAIN: C, D;	والمساورة والمساورة والمساورة والمساورة والمساورة والمساورة والمساورة والمساورة والمساورة والمساورة والمساورة	ZINC FINGER PROTEIN GLI1; CHAIN: A; DNA;	CHAIN: C, D;	•	ZINC FINGER PROTEIN	CHAIN: C, D;		ZINC FINGER PROTEIN	GLII; CHAIN: A; DNA; CHAIN: C, D;
SEQ FOLD score																94.24	
PMF score	:	0.33		0.72			1.00			0.99			0.11				
Verify		-0.26		-0.24			0.49			0.16			-0.34		٠.		
Psi Blast		3.4e-30		5.1e-32			3.46-34			3.46-34			5.1e-31			1.7e-34	
END		163		191			305			386			162			473	`.
START		2 9		06	:		174			258			26			334	:
CHAIN		ပ		ບ			Ą			¥			V			Ą	
PDB ID		lubd		lubd			2gli			2gli			2gli			2gli)
SEQ ID		410		410			410			410			410			410	

Table 5

,	 7		7			array days, etc.		
PDB annotation	FINGER, COMPLEX (DNA-BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE- FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- RINDING PROTEIN/ONA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE- FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)		CALCIUM-BINDING PROTEIN 2A9, CACY, S100A6, PRA; CALCIUM- BINDING PROTEIN, EP- HAND, S-100 PROTEIN, NMR	CALCTUM/PHOSPHOLIPID BINDING PROTEIN P11, CALPACTIN LIGHT CHAIN; S100 FAMILY, EF-HAND PROTEIN, LIGAND OF ANNEXIN II, 2 CALCTUM/PHOSPHOLIPID	METAL BINDING PROTEIN S100B, S100BETA; S100BETA; S100BETA; DIPOLAR COUPLINGS, EF. HAND, S100 2 PROTEIN, CALCIUM. BINDING PROTEIN, FOUR-HELIX BUNDLE, THREE- 3 DIMENSIONAL STRUCTURE, SOLUTION STRUCTURE	CALCIUM-BINDING PROTEIN SNTNC; CALCIUM BINDING, REGULATION, IN
Compound		ZINC FINGER PROTEIN GL11; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;		CALCYCLIN (RABBIT, CA2+); CHAIN: A, B;	S100A10; CHAIN: A, B;	S-100 PROTEIN, BETA CHAIN; CHAIN: A, B;	N-TROPONIN C; CHAIN: NULL;
SEQ FOLD score				-	70.10	87.95	84.29	
PMF score		1.00	0.90					-0.07
Verify score		0.45	0.02					0.02
Psi Blast	·	1.7e-34	1e-32		1.46-18	3.36-21	6.8e-22	1.5e-24
END		470	218	i	92	96	93	84
START AA		342	82			8	2	4
CHAIN	·	Ą	4		∀	4	V	
PDB CI		2gli	2gli		1a03	1a4p	1b4c	1blq
SEQ ID NO:		410	410		411	411	411	411

Table 5

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PDB annotation	TROPONIN C, SKELETAL MUSCLE, 2 CONTRACTION	METAL TRANSPORT CALMODULIN, HIGH RESOLUTION, DISORDER	CALCIUM-BINDING CALCIUM-BINDING, ZINC, METAL-BINDING, ACETYLATION			į	RECEPTOR RECEPTOR,	IL-6 TYPE CYTOKINES,	THIRD 2 N-TERMINAL		TRANSMEMBRANE, GLYCOPROTEIN	RECEPTOR RECEPTOR,	SIGNAL TRANSDUCER OF	TUPO LIFE CITONINES,	DOMAIN.	NE,		HORMONE/GROWTH	_	GROWTH	TIN A71, TIN; TITIN,
Compound		CALMODULIN; CHAIN: A;	S-100 PROTEIN; CHAIN: NULL;	CONTRACTILE SYSTEM PROTEIN TROPONIN C 11OP 3	MUSCLE PROTEIN TROPONIN C (TRIC FRAGMENT) (APO FORM) (NMR, 1 STRUCTURE) 1TRF 3		GP130; CHAIN: NULL;					GP130; CHAIN: NULL;			•	<u>:</u>		GROWTH HORMONE;	CHAIN: A; FROLACIAN RECEPTOR: CHAIN: B:		TITIN; CHAIN: NULL;
SEQ FOLD score		·	83.93		·		51.99				·				. •			57.27			·
PMF		-0.15		0.07	0.15							0.81									-0.14
Verify		0:00		0.21	0.26							0.28									0.00
Psi Blast		1.4e-31	1.7e-20	6.80-27	1.5e-24		3.4e-16					3.4e-16						1.26-21			5.1e-09
AA BEND		66	06	. 66	84		118					111						218			205
START AA		4	3	4	4		15					76						77			116
CHAIN		Ą																В			
208 ED ED		lexr	Imho	1top	ltf		1bj8					15j8			-			1bp3			1bpv
SEQ ID NO:		411	411	411	411		413					413						413			413

Table

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PDB annotation	CONNECTIN, FIBRONECTIN TYPE III	CONNECTIN A71, CONNECTIN; TITIN, CONNECTIN, FIBRONECTIN TYPE III	CONNECTIN A71, CONNECTIN; TITIN, CONNECTIN, FIBRONECTIN TYPE III	SIGNALING PROTEIN CYTOKINE RECEPTOR, GLYCOPROTEIN 130, GP130, INTERLEUKINE 6 2 RECEPTOR BETA SUBUNIT, SIGNALING PROTEIN	SIGNALING PROTEIN CYTOKINE RECEPTOR, GLYCOPROTEIN 130, GP130, INTERLEUKINE 6 2 RECEPTOR BETA SUBUNIT, SIGNALING PROTEIN	MEMBRANB PROTEIN BETA SANDWICH, CYTOKINE TECEPTOR, FN3 DOMAIN		ROTEIN BINDINC CYTOKINE	COMPLEX
Compound		TITIN; CHAIN: NUIL;	TITIN; CHAIN: NULL;	GP130; CHAIN: A, B;	OP130; CHAIN: A, B;	CYTOKINE RECEPTOR COMMON BETA CHAIN; CHAIN: A;	NEURAL ADHESION MOLECULE DROSOPHILA NEUROGLIAN (CHYMOTRYPTIC FRAGMENT CONTAINING THE 1CFB 3 TWO AMINO PROXIMAL FIBRONECTIN TYPE III REPEATS 1CFB 4 (RESIDUES 610 - 814)) 1CFB 5	GRANULOCYTE COLONY- STIMULATING FACTOR RECEPTOR; CHAIN: NULL;	ERYTHROPOIETIN;
SEQ FOLD score			·	52.35			64.96	·	
PMF score		0.28	0.28		0.37	0.15		0.24	0.58
Verify score		0.12	-0.10		5 0.0	0.09		-0.31	-0.05
Psi Blast		2.3e-18	1.7e-14	1.4e-16	1.26-17	1.3e-14	5.1e-24	2e-12	9.96-17
END		119	114	231	124	114	224	108	114
START AA		21	32	20	5 6	18	18	21	18
CHAIN		·		K	⋖	A			В
PDB III		1bpv	lbpv	1bgu	1bqu	1c8p	1cfb	1cto	leer
SEQ ID		413	413	413	413	413	413	413	413

Table

ation	EPTOR) OPOETIN, IN IAL 2 COKINE ASS 1, EPTOR)		PROTEIN LULAR	TTEGRIN IN AND NG	TTEGRIN IN AND NG	PROTEIN, COLUMB TO LULAR HIN.	PROTEIN, D. LULAR C. RIN- PROTEIN, C.	PROTEIN DI PROTEIN DI LULAR TU RIN.	OTEIN
PDB annotation	(CYTOKINE/RECEPTOR) EPOBP; ERYTHROPOIETIN, ERYTHROPOIETIN RECEPTOR, SIGNAL 2 TRANSDUCTION, HEMATOPOIETIC CYTOKINE, CYTOKINE RECEPTOR 3 CLASS 1, COMPLEX (CYTOKINE/RECEPTOR)		CELL ADHESION PROTEIN RGD, EXTRACELLULAR MATRIX IFNF 18	HEPARIN AND INTEGRIN BINDING HEPARIN AND INTEGRIN BINDING	HEPARIN AND INTEGRIN BINDING HEPARIN AND INTEGRIN BINDING	CELL ADHESION PROTEIN, F CELL ADHESION PROTEIN, F RGD, EXTRACELLULAR MATRIX, 2 HEPARIN- BINDING, GLYCOPROTEIN	CELL ADHESION PROTEIN, RGL, EXTRACELLULAR MATRIX, 2 HEPARIN-BINDING, GL YCOPROTEIN	CELL ADHESION PROTEIN CELL ADHESION PROTEIN, LA RGD, EXTRACELLULAR MATRIX, 2 HEPARIN-BINDING, GLYCOPROTEIN [1]	STRUCTURAL PROTEIN
Compound	CHAIN: A; ERYTHROPOIETIN RECEPTOR; CHAIN: B, C;	CELL ADHESION PROTEIN FIBRONECTIN CELL- ADHESION MODULE TYPE III-10 1FNA 3	FIBRONECTIN; 1FNF 6 CHAIN: NULL; 1FNF 7	FIBRONECTIN; CHAIN: A;	FIBRONECTIN; CHAIN: A;	FIBRONECTIN; CHAIN: NULL;	FIBRONECTIN; CHAIN: NULL;	FIBRONECTIN; CHAIN: NULL;	INTEGRIN BETA-4
SEQ FOLD score			92.46	81.90		60.33			66.57
PMF		0.69	-		0.19	٠.	0.07	0.64	
Verify		-0.27			-0.65		-0.02	-0.40	
Psi Blast		8.5e-14	8.5e-32	le-27	5.le-15	1.7e-26	9.9e-19	1.7e-14	120-25
END		107	362	293	106	207	157	107	231
START		87	21	23	E .	21	23	4	21
CHAIN			·	ď	4				
PDB CI		Ifna	1fnf	1fnh	1fnh	1mfn	1mfn	e di la companya di l	1003
SEQ ID		413	413	413	413	413	413	413	413

Table 5

			·	PU	TA	502/1	HESTI
PDB annotation	INTEGRIN, HEMIDESMOSOME, FIBRONECTIN, CARCINOMA, STRUCTURAL 2 PROTEIN	STRUCTURAL PROTEIN INTEGRIN, HEMIDESMOSOME, FIBRONECTIN, CARCINOMA, STRUCTURAL 2 PROTEIN	STRUCTURAL PROTEIN TENASCIN, FIBRONECTIN TYPE-III, HEPARIN, EXTRACELLULAR 2 MATRIX, ADHESION, FUSION PROTEIN, STRUCTURAL PROTEIN	STRUCTURAL PROTEIN TENASCIN, FIBRONECTIN TYPE-II, HEPARIN, EXTRACELLULAR 2 MATRIX, ADHESION, FUSION PROTEIN, STRUCTURAL PROTEIN		PROTEIN BINDING ED-B, FIBRONECTIN, TYPEIII DOMAIN, ANGIOGENESIS, PROTEIN 2 BINDING	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN
Compound	SUBUNIT; CHAIN: A, B;	INTEGRIN BETA-4 SUBUNIT; CHAIN: A, B;	TENASCIN; CHAIN: A, B;	TENASCIN; CHAIN: A, B;	GLYCOPROTBIN FIBRONECTIN (TENTH TYPE III MODULE) (NIMR, 36 STRUCTURES) 1TTF 3	FIBRONECTIN; CHAIN: A;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B,
SEQ FOLD score			70.56				
PMF score		0.35		0.28	0.37	0.95	0.04
Verify score		0.16		0:00	-0.29	0.29	-0.68
Psi Blast		1.2e-25	3.4e-18	3.4e-18	1.7e-14	3e-15	1.46-29
END		208	206	204	107	110	195
START		5 2	23	27	3 %	21	122
CHAIN		∢	<	<		A	V
PDB C		1983	1qr4	1qr4	1111	2fnb	laih ,
SEQ ID		413	413	413	413	413	414

	_				finds there relies	And the state of the state of	1) all may make
PDB annotation		COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN		COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN (U DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX	COMPLEX (ZINC FINGER/DNA) ZINC FINGER PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER PROTEIN-DNA INTERACTION, PROTEIN [1] DESIGN, 2 CRYSTAL
Compound	Ü	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	DNA-BINDING PROTEIN HUMAN ENHANCER- BINDING PROTEIN MBP-1 MUTANT WITH CYS 11 1BBO 3 REPLACED BY ABU (C11ABU) (NMR, 60 STRUCTURES) 1BBO 4	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;
SEQ FOLD score							
PMF score		0.01	0.25	0.46	0.21	0:00	0.41
Verify score		-0.42	0.64	0.08	-0.63	-0.18	0.37
Psi Blast		5.1e-29	9.9e-06	0.0033	6.8e-47	5.1e-38	0.0012
END		229	595	120	195	4 9	604
START		143	531	531	119	2	531
CHAIN		∢	⋖		υ	ပ	ပ
PDB D		lalh	laih	1bbo	Imey	Imey	1mey
SEQ ID		414	414	414	414	414	414

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PDB annotation	STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DINA) ZINC FINGER, PROTFIN-DINA	INTERACTION, PROTEIN	DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC FINGER, FINGER, ZINC FINGER)	PROTEIN-DNA	DESIGN, 2 CRYSTAL	STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (TRANSCRIPTION REGIT ATTONONA) YING-	YANG 1; TRANSCRIPTION	INITIATION, INITIATOR	ELEMENT, YY1, ZINC 2	PROTEIN RECOGNITION. 3	COMPLEX (TRANSCRIPTION)	REGULATION/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DNA) YING-	YANG I; IKANSCKIPTION	FIRMENT VVI ZINC 2	FINGER PROTEIN, DNA-	PROTEIN RECOGNITION, 3	COMPLEX (TRANSCRIPTION	REGULATION/DNA)	COMPLEX (IKANSCKIPTION) REGULATIONDNA) YING-	Z	INITIATION, INITIATOR ELEMBNT, YY1, ZINC2
Compound		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN: CHAIN: C. F. G:		•		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;			YY1; CHAIN: C; ADENO-ASSOCIATED VIRIS P5	INITIATOR ELEMENT	DNA; CHAIN: A, B;					YY1; CHAIN: C; ADENO-	ASSOCIATED VIRUS P5	INITIATION ELEMENT	DNA; CHAIN: A, B;		,		(A.S.A.A.A.A.A.A.A.A.A.A.A.A.A.A.A.A.A.A	ASSOCIATED VIRUS P5	INITIATOR ELEMENT	DNA; CHAIN: A, B;
SEQ FOLD score																											
PMF score		0.00				0.35			-	0.05							0.81								0.30		
Verify score		-0.33				-0.19				-0.49							-0.26				•				0.21		
Psi Blast		3.4e-13				6.8e-14				5.1e-34							3e-16								1.66-07		
END AA		195								229							193								9		
START AA		168				37				122							43								227		
CHAIN		Ð				ົນ				ပ							၁								ນ_		
PDB ED		1mey				lmey				lubd							lubd			_				,	Inpq	_	
SEQ ID NO:		414				414				414							414								414		

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PDB annotation	FINGER PROTEIN, DNA- PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)			LIGASE CBL, UBCH7, ZAP-	70, E2, UBIQUITIN, E3, PHOSPHORYLATION, 2	TYROSINE KINASE,	UBIQUITINATION, PROTEIN	DEGRADATION,	METAL BINDING PROTEIN RING FINGER PROTEIN	MATI; RING FINGER	DNA-BINDING PROTEIN	V(D)J RECOMBINATION 1	RAGI, V(D)J	RECOMBINATION,	ANTIBODY, MAD, RING	FINGER, 2 ZINC BINOCLEAR	DNA-BINDING PROTEIN	 KINASE KINASE, SIGNAL 14 TRANSDUCTION, 14 CALCTUM/CALMODULIN 11
Compound		COMPLEX(TRANSCRIPTIO N REGULATION/DNA) TRAMTRACK PROTEIN (TWO ZINC-FINGER PEPTIDE) COMPLEXED WITH 2DRP 3 DNA 2DRP 4	VRUS EQUINE HERPES VRUS-1 (C3HC4, OR RING DOMAIN) 1CHC 3 (NMR, 1 STRUCTURE) 1CHC 4	SIGNAL TRANSDUCTION	PROTEIN CBL; CHAIN: A; ZAP-70 PEPTIDE: CHAIN:	B; UBIQUITIN-	CONTUGATING ENZYME	E12-18 KDA UBCH7; CHAIN: C;	CDK-ACTIVATING KINASE ASSEMBLY	FACTOR MAT1; CHAIN: A;	RAGI; CHAIN: NULL;		٠.				•	CALCIUM/CALMODULIN- DEPENDENT PROTEIN KINASE; CHAIN: NULL;
SEQ FOLD score								•										110.80
PMF score		0.03	9.66	0.48					0.40		0.05							
Verify score		0.48	-0.12	-0.52					0.50		-0.36						_	
Psi Blast		3.3e-06	9.9e-14	1e-06	_				3.3e-13		5.1e-12							5.1e-77
END		595	61	-99					89		104					•		398
START AA		531	12	91				•	12		3							87
CHAIN		Ą		Α _				,	A									
PDB ID		2drp	Ichc	1fbv					5281		Irmd							1a06
SEQ ID NO:		414	415	415					415		415							416

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	KINASE KINASE, SIGNAL TRANSDUCTION, CALCIUM/CALMODULIN	TRANSFERASE TRANSFERASE, SERINE/THREONINE- PROTEIN KINASE, CASEIN KINASE, 2 SER/THR KINASE						PROTEIN KINASE CDK2; PROTEIN KINASE, CELL CYCLE,
Compound	CALCTUM/CALMODULIN- DEPENDENT PROTEIN KINASE; CHAIN: NULL;	PROTEIN KINASE CK2/ALPHA-SUBUNIT; CHAIN: NULL;	TRANSFERASE(PHOSPHO TRANSFERASE) \$C-/AMP\$- DEPENDENT PROTEIN KINASE (B.C.2.7.1.37) (\$C/APK\$) 1APM 3	(CATALYTIC SUBUNIT) ALPHA ISOENZYME MUTANT WITH SER 139 1APM 4 REPLACED BY	ALA (/S139A\$) COMPLEX WITH THE PEPTIDE 1APM 5 INHIBITOR PKI(5-24) AND THE DETERGENT MEGA-8 1APM 6	TRANSFERASE(PHOSPHO TRANSFERASE) \$C-/AMP\$- DEPENDENT PROTEIN KINASE (B.C.2.7.1.37) (\$C/APK\$) 1APM 3 (CATALYTIC SUBUNIT)	ALPHA ISOBNZYMB MUTANT WITH SER 139 1APM 4 REPLACED BY ALA (/S139A\$) COMPLEX WITH THE PEPTIDE 1APM 5 INHIBITOR PKI(5-24) AND THE DETERGENT MEGA-8 1APM 6	CYCLIN-DEPENDENT PROTEIN KINASE 2; CHAIN: NUIL;
SEQ FOLD score		105.92	.:			178.54	·	98.38
PMF score	1.00		96.0					·
Verify	-0.24		0.11					
Psi Blast	5.1e-77	1.2e-35	0			0		1.4e-51
END	384	400	413			410		407
START	26	99	61			89	·	94
CHAIN			ш			ជា	٠.	
PDB UD	1a06	1a60	lapm			Іарт		laq1
SEQ ID	416	416	416			416		416

				FCTA	502/	ulest 455tu
PDB annotation	PHOSPHORYLATION, STAUROSPORINB, 2 CELL DIVISION, MITOSIS, INHIBITION	COMPLEX (ISOMERASE/PROTEIN KINASE) FKBP12; SERINE/THREONINE- PROTEIN KINASE RECEPTOR R4; COMPLEX (ISOMERASE/PROTEIN KINASE), RECEPTOR 2 SERINE/THREONINE KINASE	COMPLEX (INHIBITOR PROTEIN/KINASE) INHIBITOR PROTEIN, CYCLIN-DEPENDENT KINASE, CELL CYCLE 2 CONTROL, ALPHA/BETA, COMPLEX (INHIBITOR PROTEIN/KINASE)		The state of the state of	and the latest transport
Compound		FKS06-BINDING PROTEIN; CHAIN: A, C, E, G; TGF-B SUPERFAMILY RECEPTOR TYPE I; CHAIN: B, D, F, H;	CYCLIN-DEPENDENT KINASB 6; CHAIN: A; P19INK4D; CHAIN: B;	PHOSPHOTRANSFERASE CAMP-DEPENDENT PROTEIN KINASE CATALYTIC SUBUNIT 1CMK 3 (E.C.2.7.1.37)	PHOSPHOTRANSFERASE CAMP-DEPENDENT PROTEIN KINASE CATALYTIC SUBUNIT 1CMK 3 (B.C.2.7.1.37) 1CMK 4	TRANSFERASE(PHOSPHO TRANSFERASE) CAMP- DEPENDENT PROTEIN KINASE (E.C.2.7.1.37) (CAPK) 1CTP 3
SEQ FOLD score		93.15	94.82	180.11		179.90
PMF					0.94	
Verify score					0.04	
Psi Blast		2e-25	6.8e-45	0	0	0
END		382	400	410	413	407
START		59	72	09	61	09
CHAIN		ф	₹	ш	畄	M
PDB		1960	Ibix	1cmk	1cmk	lctp
SEQ ID		416	416	416	416	416

PDB annotation									B; KINASE KINASE, TWITCHING
Compound	(CATALYTIC SUBUNIT) 1CTP 4	TRANSFERASE(PHOSPHO TRANSFERASE) CAMP- DEPENDENT PROTEIN KINASE (E.C.2.7.1.37) (CAPK) 1CTP 3 (CATALYTIC SUBUNIT) 1CTP 4	SYNTAXIN-1A; CHAIN: A, B, C;	SERINE/THREONINE- PROTEIN KINASB PAK- ALPHA; CHAIN: A, B; SERINE/THREONINE- PROTEIN KINASB PAK- ALPHA; CHAIN: C, D;	HUMAN CYCLIN- DEPENDENT KINASE 2; CHAIN: NULL;	C-JUN N-TERMINAL KINASE; CHAIN: NULL;	TWITCHIN; CHAIN: NULL;	TWITCHIN; CHAIN: A, B;	TWITCHIN; CHAIN: A, B;
SEQ FOLD score					109.49	91.45			. 142.88
PMF		1.00	0.18	1.00			1.00	1.00	
Verify		0.24	0.55	-0.06			0.13	0.21	
Psi Blast		0	9.9e-13	3.3e-63	1e-53	5.1e-39	1.26-71	5.1e-77	5.1e-77
END		381	515	316	407	446	357	382	422
START		61	398	78	76	78	75	7.1	73
CHAIN		ш	4	ပ		·		V	Ą
PDB		Ісф	lez3	1f3m	Ihcl	Ijnk	1koa	1kob	1kob
SEQ ID		416	416	416	416	416	416	416	416

Fable 5

PDB annotation	ACTIVATED PROTEIN KINASE; TRANSFERASE, MAP KINASE, SERINE/THREONINE- PROTEIN KINASE, 2 P38	KINASE RABBIT MUSCI.E PHOSPHORYLASE KINASE; GLYCOGEN METABOLISM, TRANSFERASE, SERINE/THREONINE- PROTEIN, 2 KINASE, ATP- BINDING	KINASE RABBIT MUSCLE PHOSPHORYLASE KINASE; GLYCOGEN METABOLISM, TRANSFERASE, SERINETHREONINE- PROTEIN, 2 KINASE, ATP- BINDING, CALMODULIN- BINDING	TRANSFERASE MAP KINASE, SERINE/THREONINE PROTEIN KINASE, TRANSFERASE SERINE KINASE SERINE KINASE TITIN MUSCLE.	AUTOINHIBITION SERINE KINASE SERINE KINASE, TITIN, MUSCLE,	SERINE KINASE SERINE KINASE, TITIN, MUSCLE, AUTOINHIBITION	TRANSFERASE MITOGEN FOR ACTIVATED PROTEIN FUR KINASE, MAP 2, ERK2; FUR TRANSFERASE, FUR
Compound	NUIL;	PHOSPHOR YLASB KINASE; CHAIN: NULL;	PHOSPHORYLASE KINASE; CHAIN: NULL;	ERK2; CHAIN: NULL; TITIN; CHAIN: A, B;	TITIN; CHAIN: A, B;	TITIN; CHAIN; A, B;	EXTRACELLULAR. REGULATED KINASE 2; CHAIN: NULL;
SEQ FOLD score		126.81		108.39	119.19		108.88
PMF			1.00	1.00		1.00	ŀ
Verify			0.32	0.11		0.07	
Psi Blast		1.7e-80	1.7e-80	1.2e-39 1e-57	6.6e-64	6.6e-64	5.1e-42
END		367	356	431	405	317	436
START		93	95	85 91	16	95	62
CHAIN				4	A	A	
PDB		1phk	1рћк	1pme 1tki	1tki	1tki	3erk
SEQ ID		416	416	416	416	416	416

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														. =						
PDB annotation	SERINE/THREONINE- PROTEIN KINASE, MAP KINASE, 2 ERK2	SERINE PROTEASE SERINE PROTEINASE, TRYPSIN, HYDROLASE	SERINE PROTEASE SERINE PROTEINASE, TRYPSIN, HYDROLASE	SERINE PROTEINASE TRYPSIN-LIKE SERINE	FROIEINASE, IEIKAMEK, HEPARIN, ALLERGY, 2 ASTHMA	SERINE PROTEASE	HYDROLASE, SERINE PROTEASE	SERINE PROTEASE	HYDROLASE, SEKUNE PROTEASE	E ERTING	ENZYME (PRECE), U	KALLIKREIN, SERINE	PROTEASE, PROTEIN AND MATTIRATION	COMPLEX (BLOOD	AUTOPROTHROMBIN IIA;	HYDROLASE, SERINE [1]	PROTEINASE), PLASMA CAT CHIM BINDING 2	GLYCOPROTEIN, COMPLEX	(BLOOD COAGULATION/INHIBITOR I	SERINB PROTEASE SERINB/U PROTEASE HEADER
Compound		TRYPSIN; CHAIN: A, B, C, D;	TRYPSIN; CHAIN: A, B, C, D;	BETA-TRYPTASE; CHAIN: A, B, C, D;		ALPHA TRYPSIN; CHAIN:	A, B;	ALPHA TRYPSIN; CHAIN:	A, B;	GLANDULAR KALLIKREIN-13; CHAIN:	A, B;			ACTIVATED PROTEIN C;	MAI; CHAIN; P;	•				ALPHA THROMBIN; CHAIN: A, B, F, E;
SEQ FOLD score		172.43		124.61						132.64		•		124.65						
PMR			1.00			1.00		0.99												0.87
Verify score			0.76			0.26		-0.48										······		-0.05
Psi Blast		0	0	3.4e-81		1.70-49		1.4e-47	•	1.7e-79			_	5.1e-72						3.4e-35
A E		241	241	241		142		241		241				239					,	144
START		18	19	18		19		143		14				18						20
CHAIN		V	V	¥		A		В		А				၁				٠,		В
PDB UD		1a0j	1a0j	1a01		1aks		laks		1ao5				laut						1ррх
SEQ ID		417	417	417		417		417		417				417						417

Table 5

PDB annotation	HETNAM	SERINE PROTEASE SERINE PROTEASE HEADER HETNAM	SERINE PROTEASE SERINE	COMPI WARNT BACTOR D	CATAL WITC 2 TRIAD, SELECT	REGULATION	BLOOD CLOTTING TSV-PA;	FIBRINOL YSIS,	PLASMINOGEN	ACTIVATOR, SERINE	VENOTE COMPLEX	(HYDROL ASE/INHIBITOR)	BLOOD CLOTTING	BLOOD CLOTTING TSV-PA;	FIBRINOL YSIS,	PLASMINOGEN	ACTIVATOR, SERINE	PROTEINASE, 2 SNAKE	VENOM, COMPLEX	(HYDROLASE/INHIBITOR), TO BI OOD OF OUTTING	BLOOD CLOTAING	COMPLEX (SERINE PROTEASP/INHIBITOR)	INFLAMMATION.	INHIBITOR, SPECIFICITY,	SERINE PROTEASE, 2	COMPLEX (SERINE	OR)	SERINE PROTEASE	HIDROLASE, SEKINE	, Z	SIGNAL, MULTIGENE III	ASE/HYDROLASE	
Compound		ALPHA THROMBIN; CHAIN: A, B, F, E;	COMPLEMENT FACTOR D;	CHAIN: NOLL;		_	PL ASMINOGEN	ACTIVATOR; CHAIN: A, B;	GLU-GLY-ARG-	CHLOROMETHYLKETONE	INFIBITOR; CHAIN: E, F;	•		PLASMINOGEN	ACTIVATOR; CHAIN: A, B;	GLU-GLY-ARG-	CHLOROMETHYLKETONE	INHIBITOR; CHAIN: E, F;				CATHEPSIN G; CHAIN: A;	INHIBITION SUC-VAL-PRO-	PHEP-(OPH)2; CHAIN: S;		•		TRYPSIN; CHAIN: NULL;				ENTEROPEPTIDASE;	
SEQ FOLD score			133.21				154.29						÷									126.65						165.05				136.64	
PMF score		0.55						,						1.00																			
Verify score		-0.62												69.0																			
Psi Blast		1.7e-32	1.5e-68				6.8e-87							6.8e-87								8.5e-71						5.1e-96				1c-79	
END		238	239	•			241							241								25 05 0					÷	241				239]
START AA		149	17		-		17						•	18								17		•				18	_			16	
CHAIN		E.					Ą							A																		В	
PDB UD		16ћх	1bio				Ibqy							1bqy								lcgh						1dpo				lekb	1
SEQ ID NO:		417	417				417							417								417						417				417	

[able]

		·			
INHIBITOR ENTEROKINASE, HEAVY CHAIN; ENTEROKINASE, LIGHT CHAIN; ENTEROPEPTIDASE, TRYPSINOGEN ACTIVATION, 2 HYDROLASEHYDROLASE	INHIBITOR	COMPLEX (PROTEASE/INHIBITOR) TRYPSIN, COAGULATION FACTOR XA, CHIMERA, PROTEASE, PPACK, 2 CHLOROMETHYLKETONE, COMPLEX (PROTEASE/INHIBITOR)	COMPLEX (PROTEASE/INHIBITOR) TRYPSIN, COAGULATION FACTOR XA, CHIMERA, PROTEASE, PPACK, 2 CHLOROMETHYLKETONE, COMPLEX (PROTEASE/INHIBITOR)		HYDROLASBHYDROLASB NITHERTOR CHYMOTRYPSIM HE HE HE HE HE HE HE HE HE HE HE HE HE
CHAIN: A; ENTEROPEPTIDASE; CHAIN: B; VAL-ASP-ASP- ASP-ASP-LYS PEPTIDE; CHAIN: C;	HYDROLASE (SERINE PROTEASE) PORCINE B- TRYPSIN (B.C.3.4.21.4)	COAGULATION FACTOR XA-TRYPSIN CHIMERA; CHAIN: A; D-PHE-PRO- ARG- CHLOROMETHYLKETONE (PPACK) WITH CHAIN: I;	COAGULATION FACTOR XA-TRYPSIN CHIMERA; CHAIN: A; D-PHE-PRO- ARG- CHLOROMETHYLKETONE (PPACK) WITH CHAIN: 1;	HYDROLASE (SERINE PROTEINASE) GAMMA- *CHYMOTRYPSIN *A (B.C.3.4.21.1) (\$P*H 7.0) 1GCT 3	GAMMA CHYMOTRYPSIN; CHAIN: A; GAMMA CHYMOTRYPSIN; CHAIN: B; GAMMA CHYMOTRYPSIN; CHAIN: C;
		155.00		125.17	
	0.65		1.00	• .	0.94
	-0.81		0.53		0.27
	5.1e-15	1.7e-88	1.7e-88	1.5e-78	6.8e-42
	59	241	241	241	143
	19	18	19	7	19
	V	⋖	∢	¥	Ф
	lept	1fxy	1fky	1gct	1886
	417	417	417	417	417
		CHAIN: A; ENTEROPEPTIDASE; CHAIN: B; VAL-ASP-ASP-ASP-ASP-ASP-ASP-ASP-ASP-ASP-ASP	CHAIN: A; ENTEROPETIDASE; CHAIN: B; VAL-ASP-ASP-ASP-ASP-ASP-ASP-ASP-ASP-ASP-ASP	Icpt A 19 59 5.16-15 -0.81 0.65 HYDROLASE (SERINB FRODE)	House Buy Charachest House Buy Charachest

Table 5

			· · ·		PETAS	חיותי כבודי	
PDB annotation	SERINE PROTEASE SERINE PROTEASE, HYDROLASE, MAST CELL, ANGIOTENSIN, ALPHA 2 TOLUENESULFONIC ACID			SERINE PROTEINASE SERINE PROTEINASE, GLYCOPROTEIN	LOOD NATHBITOR, FACTOR; HIBITOR, AGGF, BLOOD N, 2 PLASMA, TASE, NDING, 3	HYDROLASE MICROPLASMINOGEN, fill SERINE PROTEASE, ' ZYMOGEN, CHYMOTR YPSIN 2 FAMILY HYDROLASE	GROWTH FACTOR 7S NGF; IL GROWTH FACTOR (BETA- IL NGF), HYDROLASE - SERINE
Compound	CHYMASE; CHAIN: NULL;	COMPLEX(PROTEINASE) NHIBITOR) TRYPSIN (B.C.3.4.21.4) COMPLEXED WITH INHIBITOR FROM BITTER IMCT 3 GOURD	COMPLEX(PROTEINASEJI NHIBITOR) TRYPSIN (E.C.3.4.21.4) COMPLEXED WITH INHIBITOR FROM BITTER IMCT 3 GOURD IMCT 4	NEUROPSIN; CHAIN: A, B;	PACTOR IXA; CHAIN: C, L.; D-PHE-PRO-ARG; CHAIN: I;	PLASMINOGEN; CHAIN: A, B, C, D;	NERVE GROWTH FACTOR; CHAIN: A, B, G, X, Y, Z;
SEQ FOLD score	125.55	173.18		139.50	115.05	114.05	119.87
PMF			1.00				
Verify			0.76				
Psi Blast	6.8e-72	0	0	5.1e-84	3.4e-77	1e-81	6.8e-71
END	241	241	241	239	241	241	241
START	17	18		16.		7	22
CHAIN		4	∀	∀	υ	⋖	4
PDB	1klt	1mct	Inct	Inpm	1pfx	lqrz	lsgf
SEQ ID	417	417	417	417	417	417	417

Table

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PDB annotation	PROTEINASE 2 (GAMMA- NGF), INACTIVE SERINE PROTEINASE (ALPHA-NGF)	GROWTH FACTOR 7S NGF, GROWTH FACTOR (BETA- NGF), HYDROLASE - SERINE PROTEINASE 2 (GAMMA- NGF), INACTIVE SERINE PROTEINASE (ALPHA-NGF)	GROWTH FACTOR 7S NGF; GROWTH FACTOR (BETA- NGF), HYDROLASE - SERINE PROTEINASE 2 (GAMMA- NGF), INACTIVE SERINE PROTEINASE (ALPHA-NGF)	COMPLEX (SERINE PROTEASEINHIBITOR) TRYPSIN INHIBITOR; SERINE PROTEASE, INHIBITOR, COMPLEX, METAL BINDING SITES, 2 PROTEIN ENGINEERING, PROTEASE-SUBSTRATE INTERACTIONS, 3	COMPLEX (SERINE PROTEASE/INHIBITOR) TRYPSIN INHIBITOR; SERINE PROTEASE, INHIBITOR, COMPLEX, METAL BINDING SITES, 2 PROTEIN ENGINEERING, PROTEASE-SUBSTRATE INTERACTIONS, 3 METALLOPROTEINS	1222
Compound		NERVE GROWTH FACTOR; CHAIN: A, B, G, X, Y, Z;	NERVE GROWTH FACTOR; CHAIN: A, B, G, X, Y, Z;	ECOTIN; CHAIN: A; ANIONIC TRYPSIN; CHAIN: B;	ECOTIN; CHAIN: A; ANIONIC TRYPSIN; CHAIN: B;	HYDROLASE(SERINE PROTEINASE) TONIN (E.C. NUMBER NOT ASSIGNED) ITON 4
SEQ FOLD score		134.19		165.58		145.44
PMF score			1.00		1.00	
Verify score			0.60		0.67	
Psi Blast		1.7e-88	1.7e-88	1.2e-97	1.22-97	16-79
END AA		241	241	241	241	241
START		16	22	18	61	18
CHAIN		_ව	5	Ф	m .	
PDB		1sgf	lsgf	1slw	Islw	Iton
SEQ ID		417	417	417	417	417

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PDB annotation										-	COMPLEX (SERINE	PROTEASE/COAGULATION)	PROTEASE/COAGULATION),	SERINE, PROTEASE, 2	THROMBIN	COMPLEX (SERINE	PROTEASE/COAGULATION)	PROTEASE/COAGULATION).	SERINE, PROTEASE, 2		COMPLEX (SERINE C) PROTEASE/PEPTIDE)			PROTEASE/PEPTIDE), 2 H	/	
Compound	HYDROLASE (SERINE PROTEINASE) TRYPSIN (E.C.3.4.21.4) COMPLEXED	WITH THE INHIBITOR ITRN 3 DIISOPROPYL	FLUOROPHOSPHOFLUORI	DAIE (DET) IIM 4 HUMAN TRYPSIN, DEP INHIBITED ITRN 6	HYDROLASE (SERINE	(E.C.3.4.21.4) COMPLEXED	WITH THE INHIBITOR	TIKN 3 DIISOPKOPTL FI 110ROPHOSPHOFI 110RI	DATE (DFP) 1TRN 4	HUMAN TRYPSIN, DFP INHIBITED 1TRN 6	THROMBIN; CHAIN: L, H,	E, J, K, M, N;	AI PHA: CHAIN: F. G. I:			THROMBIN; CHAIN: L, H,	E, J, K, M, N;	AI PHA: CHAIN: F. G. I:		·	ALPHA THROMBIN;	THROMBIN; CHAIN: J, K,	M; FIBRINOPEPTIDE A-	ALPHA; CHAIN: F, N;	SERINE PROTEASE GAMMA-THROMBIN 2HNT	9
SEQ FOLD score	177.94																		·							
PMF				.*	1.00						0.58					96.0					0.59				0.05	
Verify	-		-		69.0	-					-0.44					0.11					-0.47				-0.39	-
Psi Blast	1.7e- 100				1.7e-	3					5.1e-32					3.4e-34					5.1e-32				6.8e-12	
END	241				241						238					144					238				70	
START	18				19						149					20					149				70	
CHAIN	A				A						E					н			. ——	٠.	×				C	
PDB	 				IEI IEI						lucy					lucy					1ycp				2hnt	
SEQ ID	417				417						417					417					417				417	

				1		P			
PDB annotation						YE JON, JGEN,	SERINE PROTEASE HYDROLASE, SERINE PROTEASE, DIGESTION, PANCREAS, 2 ZYMOGEN, SIGNAL		IMMUNE SYSTEM CATALYTIC ANTIBODY,
Compound	SERINE PROTEINASE KALLIKREIN A (B.C.3.4.21.8) 2PKA 4	SERINE PROTEINASE KALLIKREIN A (B.C.3.4.21.8) 2PKA 4	HYDROLASE(SERINE PROTEINASE) TRYPSIN (E.C.3.4.21.4) COMPLEXED WITH BENZAMIDINE INHIBITOR 2TBS 3	HYDROLASE(SERINE PROTEINASE) TRYPSIN (E.C.3.4.21.4) COMPLEXED WITH BENZAMIDINE INHIBITOR 2TBS 3	SERINE PROTEINASE RAT MAST CELL PROTEASE /II\$ (RMCPII\$) 3RP2 4	BETA TRYPSIN; CHAIN: NULL;	BETA TRYPSIN; CHAIN: NULL;	COMPLEX (ANTIBODY/ANTIGEN) HYHEL-5 FAB -COMPLEXED WITH BOBWHITE QUAIL LYSOZYME 1BQL 3 1BQL 95	CATALYTIC ANTIBODY 1B9 (LIGHT CHAIN);
SEQ FOLD score		·	168.64		116.86	170.27			
PMF score	0.98	1.00		1.00			1.00	0.12	0.01
Verify score	0.05	-0.25		0.65			0.84	-0.29	-0.18
Psi Blast	3.4e-25	3.4e-54	5.1e-97	5.1e-97	1.4e-69	8.5e-98	8.5e-98	3.4e-06	3.4e-05
END AA	8	241	241	240	239	241	241	197	197
START		94	18	- 19	17	18	29	34	23
CHAIN	Æ	m			¥			Ħ	Н
PDB	2pka	2pka	2tbs	2tbs	3rp2	çptp	брф	1bq1	lcle
SEQ ID	417	417	417	417	417	417	417	422	422

PDB annotation	DIELS-ALDER, IMMUNOGLOBULIN	IMMUNOGLOBULIN IMMUNOGLOBULIN, FAB COMPLEX, IDIOTOPE, ANTI-		IMMUNOGLOBULIN IMMUNOGLOBULIN, FAB COMPLEX, IDIOTOPE, ANTI-	DIOTOPE	IMMUNE SYSTEM VON WILLEBRAND FACTOR,	CLICOFROITEM DA (A:ALPHA) BINDING, 2 COMPLEX	(WILLEBRAND/IMMUNOGL'S OBULIN), BLOOD COAGULATION TYPE 3 2B = VON WILLEBRAND	502/C		
Compound	CHAIN: L; CATALYTIC ANTIBODY 1E9 (HEAVY CHAIN); CHAIN: H;	IG HEAVY CHAIN V REGIONS; CHAIN: A; IG HEAVY CHAIN V PEGIONS: CHAIN: R: IG	HEAVY CHAIN V REGIONS; CHAIN C HEAVY CHAIN V REGIONS; CHAIN: D;	IG HEAVY CHAIN V REGIONS; CHAIN: A; IG HEAVY CHAIN V	REGIONS; CHAIN: B; IG HEAVY CHAIN V REGIONS; CHAIN: C; IG HEAVY CHAIN V REGIONS; CHAIN: D;	IMMUNOGLOBULIN NMC- 4 IGG1; CHAIN: L;	IMMUNOGLOBULIN NMC- 4 IGG1; CHAIN: H; VON WILLEBRAND FACTOR;	CHAIN: A;	IMMUNOGLOBULINVIRU S HEMAGGLUTININ IGGZA FAB FRAGMENT (FAB 26/9) COMPLEXED WITH INFLUENCE	HEMAGGLUTININ HAT (STRAIN X47) (RESIDUES 101 - 108) IFRG 4	IMMUNOGLOBULIN FAB FRAGMENT OF
SEQ FOLD score	·	65.86			·				66.17		66.74
PMF score				0.42		0.01					
Verify				-0.09		0.17				·	
Psi Blast		1.4e-05		1.4e-05		1.7e-05			1e-05		5.1e-05
END		228		197	÷	197			226		228
START		21		35		35		·	23		21
CHAIN		Ω		Ω		Н			Ħ		Д
PDB		1cic		1cic		1fns			1frg		1fvd
SEQ ID		422		422		422			422		422

PDB annotation			COMPLEX (IMMUNORECEPTORIMMU NOGLOBULIN) COMPLEX (IMMUNORECEPTORIMMU NOGLOBULIN)	IMMUNOGLOBULIN FAB FRAGMENT, IMMUNOGLOBULIN	IMMUNOGLOBULIN FAB FRAGMENT, IMMUNOGLOBULIN	IMMUNB SYSTEM PRELIMINARY, IMMUNE SYSTEM	PETA	COMPLEX (IMMUNOGLOBULIN/AUTO) ANTIGEN) COMPLEX [I] (IMMUNOGLOBULIN/AUTO, ANTIGEN), RHEUMATOD FACTOR 2 AUTO-ANTIBODY COMPLEX	INSECT IMMUNITY INSECTIVE IMMUNITY, LPS-BINDING, ILL HOMOPHILIC ADHESION ILL
Compound	HUMANIZED ANTIBODY 4D5, VERSION 4 1FVD 3	IMMUNOGLOBULIN FAB D44.1 (IGG1,KAPPA) (BALB/C MOUSE, MONOCLONAL ANTIBODY) 1MLB 5	NIS ALPHA-BETA T-CELL RECEPTOR; CHAIN: A, B, C, D; H57 FAB; CHAIN: E, F, G, H	FAB1583; CHAIN: L, H	FAB1583; CHAIN: L, H	IGG3-KAPPA ANTIBODY (LIGHT CHAIN); CHAIN: A, C; IGG3-KAPPA ANTIBODY (HEAVY CHAIN); CHAIN: B, D;	IMMUNOGLOBULIN FAB FRAGMENT FROM HUMAN IMMUNOGLOBULIN IGGI (LAMBDA, HIL) 8FAB 3	IGG4 RBA; CHAIN: A; RF- AN IGM/LAMBDA; CHAIN: H, L;	HEMOLIN; CHAIN: A, B;
SEQ FOLD score				67.89			65.00		
PMF score		0.07	0.42		0.19	0.13		60.0	-0.02
Verify		-0.26	0.07		0.02	-0.34		0.08	0.12
Psi Blast		3.4e-07	1.7e-05	5.1e-07	5.1e-07	3.4e-05	0.00017	6.8e-13	1.4e-23
END		197	197	228	229	151	226	863	715
START		24	34	21	23	34	21	733	367
CHAIN		Д	ĬĽ,	н	Ħ	В	æ	1	4
PDB		Imb	Infd	Inld	Inld	1r24	8fab	ladq	1bih
SEQ ID		422	422	422	422	422	422	423	423

PDB annotation	INSECT IMMUNITY INSECT IMMUNITY, LPS-BINDING, HOMOPHILIC ADHESION	RECEPTOR RECEPTOR, SIGNAL TRANSDUCER OF	IL-6 TYPE CYTOKINES.	THIRD 2 N-TERMINAL	DOMAIN, TRANSMEMBRANE,	GLYCOPROTEIN	CONNECTIN A71,	CONNECTIN; IIIIN,	TYPE III	CELL ADHESION NEURAL CELL ADHESION	CELL ADHESION NEURAL CELL ADHESION	CELL ADHESION NEURAL CELL ADHESION	CELL ADHESION NEURAL CELL ADHESION	GROWTH FACTOR/GROWTH	FGFR, IMMUNOGLOBULIN	LIKE, SIGNAL TRANSDUCTION. 2	DIMERIZATION, GROWTH	FACTOR/GROWTH FACTOR	11	FACTOR RECEPTOR FOFF, [1]	LIKE, SIGNAL	TRANSDUCTION, 2	DIMERIZATION, GROWIH FACTORIL	RECEPTOR	GROWTH FACTOR/GROWI的
Compound	.;	GP130; CHAIN: NULL; RE	月	百?	200		TITIN; CHAIN: NULL;	38	<u> </u>	AXONIN-1; CHAIN: A; CE	AXONIN-1; CHAIN: A; CE	AXONIN-1; CHAIN: A; CE	AXONIN-1; CHAIN: A; CE	-	~~~	FACTOR RECEPTOR 1; LII		FA	FIBROBLAST GROWTH GR	FACTOR 2; CHAIN: A, B; FA		CHAIN: C, D;		RE	FIBROBLAST GROWTH GR
SEQ FOLD score									·																
PMF score	0.24	9.0 20.0					0.51			0.04	0.27	0.13	0.33	0.01					-0.11						0.36
Verify	0.32	0.20					-0.07			-0.14	-0.00	-0.03	90.0	-0.09	,	-			0.00	<u></u>					0.05
Psi Blast	1.7e-29	2e-11					3e-12			1.7e-21	1.7e-34	5.1e-27	1.5e-31	8.5e-20					8.5e-11					,	8.5e-26
END	888	541					528			628	722	814	888	536					725						888
START	556	442					442	_		300	351	441	120	354					240	2					729
CHAIN	4			-						A	A	A	V	ပ					C)					၁
PDB	1bih	1bj8		<u> </u>			1bpv			1cs6	1cs6	Ics6	1cs6	lcvs					Tove						lcvs
SEQ ID	423	423				•	423			423	423	423	423	423					423	}					423

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—Т				PCT/850		1 1 1 1	
PDB annotation	FACTOR RECEPTOR FGF, FGFR, IMMUNOGLOBULIN- LIKE, SIGNAL TRANSDUCTION, 2 DIMERIZATION, GROWTH FACTOR/GROWTH FACTOR RECEPTOR	GROWTH FACTOR/GROWTH FACTOR RECEPTOR FGF, FGFR, IMMUNOGLOBULIN- LIKB, SIGNAL TRANSDUCTION, 2 DIMERIZATION, GROWTH FACTOR/GROWTH FACTOR RECEPTOR	GROWTH FACTOR/GROWTH FACTOR RECEPTOR FGF, FGFR, IMMUNOGLOBULIN- LIKE, SIGNAL TRANSDUCTION, 2 DIMERIZATION, GROWTH FACTOR/GROWTH FACTOR		TA T	Ą	CELL ADHESION NCAM; I'L
Compound	FACTOR 2; CHAIN: A, B; FIBROBLAST GROWTH FACTOR RECEPTOR 1; CHAIN: C, D;	FIBROBLAST GROWTH FACTOR 2; CHAIN: A, B; FIBROBLAST GROWTH FACTOR RECEPTOR 1; CHAIN: C, D;	FIBROBLAST GROWTH FACTOR 2; CHAIN: A, B; FIBROBLAST GROWTH FACTOR RECEPTOR 1; CHAIN: C, D;	IMMUNOGLOBULIN FAB' FRAGMENT OF THE DB3 ANTI-STEROID MONOCLONAL ANTIBODY 1DBB 3 (IGG1, SUBGROUP 2A, KAPPA 1) COMPLEX WITH PROGESTERONE 1DBB 4	NEURAL CELL ADHESION MOLECULE; CHAIN: A, B, C, D;	NEURAL CELL ADHESION MOLECULE; CHAIN: A, B, C, D;	NEURAL CELL ADHESION MOLECULE; CHAIN: A, B,
SEQ FOLD score							
PMF		-0.07	0.70	0.10	0.09	0.10	-0.07
Verify		0.19	0.26	-0.31	0.19	0.22	0.02
Psi Blast		1.7e-25	6.8e-25	1.7e-06	5.1e-17	1.7e-11	1.4e-13
END		813	88	477	220	709	801
START		629	972	363	359	550	643
CHAIN		Q	Ω	н	Ą	4	Ą
PDB		Icvs	1cvs	1dbb	lepf	lepf	lepf
SEQ ID		423	423	423	423	423	423

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PDB annotation	FOLD, GLYCOPROTEIN	CELL ADHESION NCAM; NCAM, IMMUNOGLOBULIN FOLD, GLYCOPROTEIN	GROWTH PACTOR/GROWTH FACTOR RECEPTOR FGF2;	FGFR2; IMMUNOGLOBULIN	BELONGING TO THE I-SET 2-	DOMAINS, B-TREPOIL FOLD	GROWTH FACTOR/GROWTH PACTOR RECEPTOR FGF2;	FGFR2; IMMUNOGLOBULIN	(IG)LIKE DOMAINS BET ONGING TO THE I-SET 2	SUBGROUP WITHIN IG-LIKE	DOMAINS, B-TREFOIL FOLD	GROWTH FACTOR/GROWTH PACTOR RECEPTOR FGF2;	FGFR2; IMMUNOGLOBULIN	(IG)LIKE DOMAINS	BELONGING TO THE L-SEI 2	DOMAINS, B-TREFOIL FOLISH	GROWTH FACTOR/GROWTH,	FACTOR RECEPTOR FOFI:	(IG) LIKE DOMAINS	BELONGING TO THE I-SET (1)	SUBGROUP WITHIN IG-LIKE	CROWNINS, B-I KEFOLL FOLKE	FACTOR RECEPTOR FOF1;	DBULD	(IG) LIKE DOMAINS	SUBGROUP WITHIN IG-LIKE	DOMAINS, B-TREFOIL FOLD
Compound	C, D;	NEURAL CELL ADHESION MOLECULE; CHAIN: A, B, C, D;	FIBROBLAST GROWTH FACTOR 2; CHAIN: A, B, C,	D; FIBROBLAST GROWTH	CHAIN: E, F, G, H;		FIBROBLAST GROWTH	D; FIBROBLAST GROWTH	FACTOR RECEPTOR 2;	CHAME E, F, G, II,		FIBROBLAST GROWTH	D; FIBROBLAST GROWTH	FACTOR RECEPTOR 2;	CHAIN: E, F, G, H;		HBROBLAST GROWTH	FACTOR 1; CHAIN: A, B;	HISKOBLASI GROWIN	CHAIN: C, D;		THE PLANT OF THE PARTY OF THE PARTY.	FIBROBLAST GROW I H FACTOR 1; CHAIN: A, B;	FIBROBLAST GROWTH	FACTOR RECEPTOR 1;	CHAIN: C, D;	
SEQ FOLD score																								,	-	-1	
PMF		0.19	0.03				0.00	• •				0.27					0.21						60.0-				
Verify		0.11	-0.09		·	• .	0.05					-0.00					-0.07		•				0.12				
Psi		5.1e-20	1.2e-22				1.7e-23					1.4e-23					5.le-18						1.7e-10				
END		888	813				817					888					536						725				
START		731	646				646				-	739					354						250				
CHAIN		¥.	E				G					C					C						၁				
PDB	3	lepf	lev2	,			lev2					lev2					1evt	:					levt				
SEQ ID	2	423	423				423					423					423	<u> </u>					423			.	

				PLT	502/6	ILZ	
FDB annotation	GROWTH FACTOR/GROWTH FACTOR RECEPTOR FGF1; FGFR1; IMMUNOGLOBULIN (IG) LIKE DOMAINS BELONGING TO THE I-SET 2 SUBGROUP WITHIN IG-LIKE DOMAINS, B-TREFOIL FOLD	IMMUNE SYSTEM FC- EPSILON RI-ALPHA; IMMUNOGLOBULIN FOLD, GLYCOPROTEIN, RECEPTOR, IGE-BINDING 2 PROTEIN	AFFINITY IGE-FC RECEPTOR, FC(EPSILON) IGE-FC; IMMUNOGLOBULIN FOLD, GLYCOPROTEIN, RECEPTOR, IGE-BINDING 2 PROTEIN, IGE ANTIBODY, IGE-FC	AFFINITY IGE-FC RECEPTOR, FC(EPSILON) IGE-FC, IMMUNOGLOBULDE FOLD, GLYCOPROTEIN, RECEPTOR, IGE-BINDING 2 PROTEIN, IGE ANTIBODY, IGE-FC	MEMBRANE PROTEIN CD323 FC RECEPTOR, IMMUNOGLOULN, LEUKOCYTE, CD32	CONTRACTILE PROJEIN TO INMUNOGLOBULIN FOLD HE BETA BARREL	
Compound	FIBROBLAST GROWTH FACTOR 1; CHAIN: A, B; FIBROBLAST GROWTH FACTOR RECEPTOR 1; CHAIN: C, D;	HIGH AFFINITY IMMUNOGLOBULIN EPSILON RECEPTOR CHAIN: A;	HIGH APPLINITY IMMUNOGLOBULIN EPSILON RECEPTOR CHAIN: A; IG EPSILON CHAIN C REGION; CHAIN: B, D;	HIGH AFFINITY IMMUNOGLOBULIN EPSILON RECEPTOR CHAIN: A; IG EPSILON CHAIN C REGION; CHAIN: B, D;	FC RECEPTOR FC(GAMMA)RIIA; CHAIN: A;	TELOKIN; CHAIN: A	TELOKIN; CHAIN: A
SEQ FOLD score							
PMF	0.18	0.36	0.63	0.18	-0.12	0.84	0.75
Verify	0.15	0.09	90.06	0.06	-0.00	0.31	0.39
Psi	8.5e-24	3.3e-13	6.6e-12	6.6e-15	9.96-12	5.1e-16	1e-09
END	888	888	816	888	802	440	629
START	729	742	<i>1</i> 99	737	299	351	538
CHAIN	U	V	V	∢	V	Ą	4
PDB	lew	1£q	1f6a	1f6a	lfcg	1fhg	1fhg
SEQ ID	423	423	423	423	423	423	423

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PDB annotation	BETA BARREL	COMPLEX (IMMUNOGLOBULIN/RECEP	TOR) IMMUNOGLOBULIN FOLD, TRANSMEMBRANE, GI YCOPROTEIN	RECEPTOR, 2 SIGNAL,	(IMMUNOGLOBULIN/RECEP TOR)	COMPLEX (IMMUNOGLOBULIN/RECEP TOR) IMMUNOGLOBULIN	FOLD, TRANSMEMBRANE, GLYCOPROTEIN,	RECEPTOR, 2 SIGNAL,	(IMMUNOGLOBULIN/RECEPTOR)	COMPLEX (IMMUNOGLOBULIN/RECEP TOR) IMMUNOGLOBULIN		NAL,	(IMMUNOGLOBULIN/RECEPT) TOR)	KINASE KINASE, TWITCHIN INTRASTERIC REGULATION	the there are	MISCUE PROTRIN	TMS;	TRANSMEMBRANE,
Compound		INTERLEUKIN-1 BETA; CHAIN: A: TYPE 1	INTERLEUKIN-1 RECEPTOR; CHAIN: B;			INTERLEUKIN-1 BETA; CHAIN: A; TYPE 1 INTER! BITKIN-1	RECEPTOR; CHAIN: B;			INTERLEUKIN-1 BETA; CHAIN: A; TYPE 1 INTER! EIKIN-1	RECEPTOR; CHAIN: B;	<u>.</u>		TWITCHIN; CHAIN: NULL;	IMMUNOGLOBULIN IMMUNOGLOBULIN GI (IGGI) (MCG) WITH A	HINGE DELETION 1MCO 3	TITIES, CEPAINS NOTES,	
SEQ FOLD	PLUIC														83.10			
PMF	score	60.0				0.27				69:0				0.37			0.28	
Verify	Score	-0.22		:		0.07		-	:	0.19				-0.15			0.19	
Psi	Blast	3.3e-15			·.	3.3e-24				3.3e-15				1.5e-12	1.2e-11		6.8e-14	
END	¥	813				698				888				430	818		429	
START	AA	574				662				742				351	430		353	
CHAIN	e	В				В				æ					æ			
PDB	A	1itb				1itb		· -		litb				1koa	1mco		Inct	
SEQ ID	ÖN	423		·		423				423				423	423		423	

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				774 Ann 1870		
PDB annotation	REPEAT, BRAIN, 2 IMMUNOGLOBULIN FOLD, ALTERNATIVE SPLICING, SIGNAL, 3 MUSCLE PROTEIN	MUSCLE PROTEIN CONNECTIN, NEXTM5; CELL ADHESION, GLYCOPROTEIN, TRANSMEMBRANE, REPEAT, BRAIN, 2 IMMUNOGLOBULIN FOLD, ALTERNATIVE SPLICING, SIGNAL, 3 MUSCLE PROTEIN	IMMONOGLOBOLLA LLAND DOMAIN CONNECTIN 127, TITIN 1G REPEAT 27; MUSCLE PROTEIN, IMMUNOGLOBULIN-LIKE DOMAIN	IMMUNOGLOBULIN-LIKE DOMAIN CONNECTIN 127, TITIN IG REPEAT 27; MUSCLE PROTEIN, IMMUNOGLOBULIN-LIKE DOMAIN	DOMAIN CONNECTIN 127, II TITIN IG REPEAT 27; III MUSCLE PROTEIN, IMMUNOGLOBULIN-LIKE	
Compound		TITIN, CHAIN: NULL;	TITIN, 127; CHAIN: NULL;	TITIN, 127; CHAIN: NULL;	TITIN, 127; CHAIN: NULL;	MUSCLE PROTEIN TITIN MODULE M5 (CONNECTIN) 1TNM 3 (NMR, MINIMIZED AVERAGE STRUCTURE) 1TNM 4 1TNM 58
SEQ FOLD						
PMF		0.19	66'0	0.95	0.40	0.52
Verify		0.13	0.43	0.34	0.17	-0.02
Psi	Sand Sand	6.66-10	5.1e-09	9.9e-11	6.6e-12	6.8e-14
END	Y.	227	437	435	227	429
START	&	979	353	357	44	353
CHAIN	a					
PDB	a	Inct	ltit	1tit	1tit	1tmm
SEQ ID	SOS.	423	423	423	423	423

Table 5

					PETA	- 11°11	e ne ni In	
PDB annotation		CELL ADHESION PROTEIN VCAM-DI.2; IVCA 6 IMMUNOGLOBULIN SUPERFAMILY, INTEGRIN-BINDING IVCA 15	GLYCOPROTEIN CD4; INMUNOGLOBULIN FOLD, TRANSMEMBRANE, GLYCOPROTEIN, T-CELL, 2 MHC LIPOPROTEIN, POLYMORPHISM	L RAL	AL 2	IMMUNE SYSTEM CD32; # RECEPTOR, FC, CD32, IMMUNE SYSTEM	RECEPTOR, FC, CD32, IMMUNE SYSTEM IMMUNE SYSTEM	CELL ADRESION NOAM DOMAIN 1; CELL ADRESION, GLYCOPROTEIN, HEPARIN
Compound	MUSCLE PROTEIN TITIN MODULA MS (CONNECTIN) 1TNM 3 (NMR, MINIMIZED AVERAGE STRUCTURE) 1TNM 4 1TNM 58	HUMAN VASCULAR CELL ADHESION MOLECULE-1; 1VCA 4 CHAIN: A, B; 1VCA 5	T-CELL SURFACE GLYCOPROTEIN CD4; CHAIN: A, B;	MHC CLASS I NK CELL RECEPTOR PRECURSOR; CHAIN: A;	MHC CLASS I NK CELL RECEPTOR PRECURSOR; CHAIN: A;	FC GAMMA RIIB; CHAIN: A;	FC GAMMA RIIB; CHAIN: A;	NEURAL CELL ADHESION MOLECULE; CHAIN: NULL;
SEQ FOLD score		·			•		·	÷
PMF	0.33	0.42	0.86	-0.08	0.01	0.09	0.01	0.68
Verify	0.01	0.15	0.20	0.12	0.08	0.10	0.04	-0.15
Psi	6.6e-09	9.9e-11	6.6e-13	6.6e-12	2.6e-15	9.9e-16	1.3e-15	1.7e-08
END	227	802	888	803	888	816	888	430
START	199	645	737	637	741	199	737	359
CHAIN	3	₹	∢	4	4	Ą	A	
PDB		Ivca	1wio	2dli	2dli	2fcb	2fcb	2псш
SEQ ID	423 	423	423	423	423	423	423	423

Table 5

				_		Jest	there wither.	designation of the			コ
PDB annotation	BINDING, GPI-ANCHOR, 2 NEURAL ADHESION MOLECULE, IMMUNOGLOBULIN FOLD, SIGNAL				COMPLEX (BLOOD COAGULATION/INHIBITOR) AUTOPROTHROMBIN IIA; HYDROLASE, SERINE PROTEINASE), PLASMA CALCIUM BINDING, 2 GLCTUM BINDING, 2	(BLOOD COAGULATION/INHIBITOR)	HYDROLASE INHIBITOR FALL-BETA STRUCTURE, HYDROLASE INHIBITOR	SUGAR BINDING PROTEIN, UDA; LECTIN, HEVEIN (I) DOMAIN, UDA, SUPERANTIGEN,	OTEIN IN, IALLING		GLYCOPROTEIN
Compound		IMMUNOGLOBULIN IMMUNOGLOBULIN FAB' NEW (LAMBDA LIGHT CHAIN) 7FAB 3	IMMUNOGLOBULIN FAB FRAGMENT FROM HUMAN IMMUNOGLOBULIN IGGI (LAMBDA, HIL) 8FAB 3		ACTIVATED PROTEIN C; CHAIN: C, L; D-PHE-PRO- MAI; CHAIN: P;		BOWMAN-BIRK TRYPSIN INHIBITOR; CHAIN: A	AGGLUTININ ISOLECTIN VAGGLUTININ ISOLECTIN V/ CHAIN: A;	TUMOR NECROSIS PACTOR RECEPTOR; CHAIN: A, B;	LAMININ; CHAIN: NULL;	LAMININ; CHAIN: NULL;
SEQ FOLD score	·				50.32	,			58.93	72.24	
PMF		0.05	0.12				0.53	0.16			-0.08
Verify		-0.02	-0.11				-0.05	-0.43			0.09
Psi		1.7e-13	6.8e-15		3.3e-06		3.3e-17	6.6e-07	1.6e-11	6.66-15	6.6e-15
END	5	863	863		206		136	221	250	185	226
START		733	733 _		113	·	23	136	88	77	30
CHAIN	3	ы	4		<u>.</u>		¥	4	A		
PDB	3	7fab	8fab		laut		1c2a	1en2	lext	1klo	1klo
SEQ ID	Ö	423	423		424		424	424	424	424	424

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PDB annotation	GLYCOPROTEIN				KINASE KINASE, SIGNAL TRANSDUCTION, CALCIUM/CALMODULIN	KINASE KINASE, SIGNAL TRANSDUCTION, CALCIUM/CALMODULIN	TRANSFERASE TRANSFERASE, SERINE/THREONINE- PROTEIN KINASE, CASEIN KINASE, 2 SER/THR KINASE	PCT/Soe/O1	
Compound		METALLOTHIONEIN METALLOTHIONEIN ISOFORM II 4MT2 3	LECTIN (AGGLUTININ) WHEAT GERM AGGLUTININ (ISOLECTIN 2) 9WGA 3		CALCIUM/CALMODULIN- DEPENDENT PROTEIN KINASE; CHAIN: NULL;	CALCHUMICALMODULIN- DEPENDENT PROTEIN KINASE; CHAIN: NULL;	PROTEIN KINASE CK2/ALPHA-SUBUNIT; CHAIN: NULL;	TRANSFERASE (PHOSPHO TRANSFERASE) \$C-/AMP\$- DEPENDENT PROTEIN KINASE (E.C.2.7.1.37) (\$C/APK\$) 1APM 3 (CATALYTIC SUBUNIT) ALPHA ISOENZYME MUTANT WITH SER 139 1APM 4 REPLACED BY ALA (/S139A\$) COMPLEX WITH THE PEPTIDE 1APM 5 INHIBITOR PKI(5-24) AND THE DETERGENT MEGA-8 1APM 6 TRANSFERASE (PHOSPHO TRANSFERASE (PHOSPHO TRANSFERASE (PHOSPHO TRANSFERASE (PHOSPHO TRANSFERASE (PHOSPHO TRANSFERASE (PHOSPHO	DEPENDENT PROTEIN KINASE (B.C.2.7.1.37) (\$C/APK\$) 1APM 3
SEQ FOLD	21000	52.76	85.55		75.71		73.63	78.39	
PMF	PUNC					0.16	·	0.92	
Verify	SCOTE					-0.14		0.33	
Psi	Diast	1.7e-08	1.7e-18		8.5e-72	8.5e-72	6.8e-40	3.46-75	·
END	AA	185	185		394	393	401	401	· · · · · ·
START	¥¥	120	11		116	49	35	35	
Z	a		A					时	,
PDB	A	4mt2	9wga		1a06	1a06	1a60	lapm	
SEQ ID	NO:	424	424		425	425	425	425	

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PDB annotation		PROTEIN KINASSE CDKZ; PROTEIN KINASSE, CELL CYCLE, PHOSPHORYLATION, STAUROSPORINE, 2 CELL DIVISION, MITOSIS, INHIBITION	PROTEIN KINASE CDK2; PROTEIN KINASE, CELL CYCLE, PHOSPHORYLATION, STAUROSPORINE, 2 CELL DIVISION, MITOSIS, INHIBITION	COMPLEX (ISOMERASE/PROTEIN KINASE) FKBP12; SERINE/THREONINE- PROTEIN KINASE RECEPTOR R4; COMPLEX (ISOMERASE/PROTEIN KINASE), RECEPTOR 2 SERINE/THREONINE KINASE	COMPLEX (KINASE/INHIBITOR) CDK6 P19INK4D; CYCLIN DEPENDENT KINASE, [1] CYCLIN DEPENDENT [1] KINASE INHIBITORY 2 [1]
Compound	(CATALYTIC SUBUNIT) ALPHA ISOENZYME MUTANT WITH SER 139 1APM 4 REPLACED BY ALA (/S139AS) COMPLEX WITH THE PEPTIDE 1APM 5 INHIBITOR PKI(5-24) AND THE DETERGENT MEGA-8 1APM 6	CYCLIN-DEPENDENT PROTEIN KINASE 2; CHAIN: NULL;	CYCLIN-DEPENDENT PROTEIN KINASE 2; CHAIN: NUIL;	FK506-BINDING PROTEIN; CHAIN: A, C, B, G, TGF-B SUPERFAMILY RECEPTOR TYPE I; CHAIN: B, D, F, H;	CYCLIN-DEPENDENT KINASE 6; CHAIN: A, C; CYCLIN-DEPENDENT KINASE INHIBITOR; CHAIN: B, D;
SEQ FOLD score			78.41	80.91	87.11
PMF score		0.89			
Verify		-0.04		·	
Psi		1.7e-70	1.7e-70	6.66-31	6.8e-51
END		395	401	401	393
START		99	99	47	123
CHAIN	1			Ф	4
PDB	3	laq1	laq1	1b6c	16i8
SEQ ID		425	425	425	425

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PDB annotation	PROTEIN, CDK, INK4, CELL CYCLE, COMPLEX (KINASE/INHIBITOR) HBADER HELLX	COMPLEX (INHIBITOR PROTEIN/KINASE) INHIBITOR PROTEIN, CYCLIN-DEPENDENT KINASE, CELL CYCLE 2 CONTROL, ALPHABETA, COMPLEX (INHIBITOR PROTEIN/KINASE)	PHOSPHOTRANSFERASE PROTEIN KINASE 1CKI 18		-	PHOSPHOTRANSFERASE		U willia (Ema Eliza, Eliza,
Compound		CYCLIN-DEPENDENT KINASE 6; CHAIN: A; P19INK4D; CHAIN: B;	CASEIN KINASE I DELTA; ICKI 6 CHAIN: A, B; ICKI 7	PHOSPHOTRANSFERASE CAMP-DEPENDENT PROTEIN KINASE CATALYTIC SUBUNIT ICMK 3 (B.C.2.7.1.37)	PHOSPHOTRANSFERASE CAMP-DEPENDENT PROTEIN KINASE CATALYTIC SUBUNIT 1CMK 3 (B.C.2.7.1.37)	CASEIN KINASE-1; 1CSN 4	TRANSFERASE (PHOSPHO TRANSFERASE) CAMP- DEPENDENT PROTEIN KINASE (B.C.2.7.1.37) (CAPK) 1CTP 3 (CATALYTIC SUBUNIT) 1CTP 4	TRANSFERASE(PHOSPHO TRANSFERASE) CAMP- DEPENDENT PROTEIN KINASE (E.C.2.7.1.37) (CAPK) 1CTP 3
SEQ FOLD score		101.24	66.20	82.38		62.13	90.28	
PMF			. *	·	0.95			0.94
Verify					0.19			0.18
Psi		3.4e-53	1.6e-34	1.7e-76	1.7e-76	3.4e-15	1.7e-76	1.7e-76
END		401	398	401	393	393	401	393
START		8	62	3 6	50	63	32	50
CHAIN		∢	4	ш	ъ		Ħ	四
PDB		1blx	1cki	1cmk	1cmk	1csn	1ctp	lctp
SEQ ID		425	425	425	425	425	425	425

			-	Torke street, begins of	in the same of the last	
PDB annotation		TRANSFERASE KINASE DOMAIN, AUTOINHIBITORY FRAGMENT, HOMODIMER	PHOSPHOTRANSFERASE FGFR IK, FIBROBLAST GROWTH FACTOR RECEPTOR 1; TRANSFERASE, TYROSINE- PROTEIN KINASE, ATF- BINDING, 2 PHOSPHORYLATION, RECEPTOR, PHOSPHOTRANSFERASE	PHOSPHOTRANSFERASE FGERIK, FIBROBLAST GROWTH FACTOR RECEPTOR 1; TRANSFERASE, TYROSINE- PROTEIN KINASE, ATP- BINDING, 2 PHOSPHORYLATION, RECEPTOR, PHOSPHOTRANSFERASE	PROTEIN KINASE CUKZ; TRANSFERASE, SERINE/THREONINE PROTEIN KINASE, ATP- BINDING, 2 CELL CYCLE, CELL DIVISION, MITOSIS, PHOSPHORYLATION	TRANSFERASE, MEANSFERASE, MEANSFERASE, MEANSFERASE, MEANFORME MEANFORME MEANFORMER, ATP.
Compound	(CATALYTIC SUBUNIT) 1CTP 4	SERINETHREONINE- PROTEIN KINASE PAK- ALPHA; CHAIN: A, B; SERINETHREONINE- PROTEIN KINASE PAK- ALPHA; CHAIN: C, D;	FGF RECEPTOR 1; CHAIN: A, B;	FGF RECEPTOR 1; CHAIN: A, B;	HUMAN CYCLIN- DEPENDENT KINASE 2; CHAIN: NULL;	HUMAN CYCLIN- DEPENDENT KINASE 2; CHAIN: NUIL;
SEQ FOLD score			84.95	82.99		97.32
PMF	٠	0.06			0.92	
Verify		-0.05			0.09	
Pst		5.1e-68	6.6e-31	3.4e-30	8.5e-75	8.5e-75
END		393	401	401	395	401
START	e e	49	,	45	99	99
CHAIN	3	U	4	æ		
PDB	3	1f3m	1fgk	1fgk	Ihcl	1hci
SEQ ID	i)	425	425	425	425	425

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			<u> </u>			IP"tz iP"		
PDB annotation	BINDING, 2 CELL CYCLE, CELL DIVISION, MITOSIS, PHOSPHORYLATION	SERINE/THREONINE- PROTEIN KINASE CSBP, RK, P38; PROTEIN SER/THR- KINASE, SERINE/THREONINE-	COMPLEX (TRANSFERASE/SUBSTRAT E) TYROSINE KINASE, SIGNAL TRANSDUCTION, PHOSPHOTRANSFERASE, 2 COMPLEX (KINASE/PEPTIDE SUBSTRATE/ATP ANALOG), ENZYME, 3 COMPLEX (TRANSFERASE/SUBSTRAT E)	KINASE KINASE, TWITCHIN, INTRASTERIC REGULATION	KINASE KINASE, TWITCHIN, INTRASTERIC REGULATION	KINASE KINASE, TWITCHING INTRASTERIC REGULATION	TRANSFERASE MITOGEN ACTIVATED PROTEIN KINASE, TRANSFERASE, MAP KINASE, SERINE/THREONINE-	KINASE RABBIT MUSCLE IJ PHOSPHORYLASE KINASE;., GLYCOGEN METABOLISM, TRANSFERASE, SERINE/THREONINE. PROTEIN, 2 KINASE, ATP. IJ BINDING, CALMODULIN- IJ BINDING
Compound		P38 MAP KINASB; CHÁIN: NULL;	INSULIN RECEPTOR; CHAIN: A; PEPTIDE SUBSTRATE; CHAIN: B;	TWITCHIN; CHAIN: NULL;	TWITCHIN; CHAIN: A, B;	TWITCHIN; CHAIN: A, B;	MAP KINASE P38; CHAIN: NULL;	PHOSPHORYLASE KINASE; CHAIN: NULL;
SEQ FOLD score		66.27	77.78		71.13		76.87	75.55
PMF score				0.94		0.76		
Verify score				0.19		0.24		
Psi Blast		6.8e-43	9.96-30	1.7e-56	1e-58	le-58	3.4e-50	3.4e-67
END AA		401	393	393	400	393	401	378
START		50	54	99	43	59	05	92
CHAIN		·	A		Ą	Ą		
PDB		lian	11:13	1koa	1kob	1kob	1p38	1phk
SEQ ID		425	425	425	425	425	425	425

PDB annotation	KINASE RABBIT MUSCLE PHOSPHORYLASE KINASE; GLYCOGEN METABOLISM, TRANSFERASE, SERNETHREONINE- PROTEIN, 2 KINASE, ATP- BINDING, CALMODULIN- BINDING	TRANSFERASE MAP KINASE, SERINE/THREONINE PROTEIN KINASE, TRANSFERASE	SERINE KINASE SERINE KINASE, TITIN, MUSCLE, AUTOINHIBITION	TRANSFERASE MITOGEN ACTIVATED PROTEIN KINASE, MAP 2, ERK2; TRANSFERASE, SERINE/THREONINE- PROTEIN KINASE, MAP KINASE, 2 ERK2	<u> </u>	IMMUNOGLOBULIN, IN INVARIANT	5 502/(COMPLEX (IMMUNORECEPTOR/IMMUM) NOGLOBULIN) COMPLEX [U] (IMMUNORECEPTOR/IMMUM) NOGLOBULIN)
Compound	PHOSPHORYLASE KINASE; CHAIN: NULL;	ERK2; CHAIN: NULL;	TITIN; CHAIN: A, B;	EXTRACELLULAR REGULATED KINASE 2; CHAIN: NULL;		MONOCLONAL ANTIBODY D1.3; CHAIN: L, H;	IMMUNOGLOBULIN IMMUNOGLOBULIN VL DOMAIN (VARIABLE DOMAIN OF KAPPA LIGHT IIVL 3 CHAIN) OF DESIGNED ANTIBODY M29B IIVL 4	NIS ALPHA-BETA T-CELL RECEPTOR; CHAIN: A, B, C, D; H57 FAB; CHAIN: E, F, G, H
SEQ FOLD score		70.74	57.34	71.41		61.78	62.21	88.42
PMF	0.93							
Verify	0.19							
Psi	3.4e-67	1.7e-49	1.2e-46	3.46-50		5.1e-29	5.1e-28	3.4e-51
END	395	401	401	401		130	131	196
START	19	09	63	55		20	02	20
CHAIN			V V	·		T	V	æ
	1pk	Ірте	1tki	3erk		la7q	livl	Infd
SEQ ID	425 425	425	425	425		426	426	426

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PDB annotation			HYDROLASE MACROPAIN	SUBUNIT Y7, PROTEINASE	MACBOBAN STIBTINET VI3	PROTEINASE VSCE	SUBUNIT 13, MACROPAIN	SUBUNIT PRE6,	PROTEINASE YSCE	SUBUNIT MACROPAIN	SUBUNIT PUP2,	PROTEINASE YSCE	SUBUNIT MACROPAIN	SUBUNIT PRES,	PROTEINASE YSCE	SUBUNII MACKOPAIN	SUBUNIT CI, PROTEINASE	YSCE SUBUNIT 1,	MACROPAIN SUBUNIT C7-	ALPHA, PROTEINASE YSCE	MACROPAIN SUBUNIT	3	SUBUINI MACKUFALIN	SUBUNII PUPS,	MACBOBAIN STRINGT OFF	DECTAIN SOCIAL CITY	STIBITUTE 11 MACROPAIN	SUBLINIT PREZ.	PROTEINASE YSCE	SUBUNIT	MULTICATALYTIC 🔭	ENDOPEPTIDASE COMPLEX	SUBUNIT CS; MACROPAIN [1]	#
Compound	IMMUNOGLOBULIN BENCE-*JONES PROTEIN (LAMBDA, VARIABLE DOMAIN) 2RHE 4	,	PROTEASOME	COMPONENT Y7; CHAIN:	A, O; PROTEASOME	P. P. PPOTTE A COMP.	COMPONENT PRE6:	CHAIN: C, O;	PROTEASOME	COMPONENT PUP2;	CHAIN: D, R;	PROTEASOME	COMPONENT PRE5;	CHAIN: B, S;	PROTEASOME	COMPONENT C1; CHAIN:	F, T; PROTEASOME	COMPONENT C7-ALPHA;	CHAIN: G, U;	PROTEASOME	COMPONENT PUPI;	CHAIN: H, V;	PROTEASOME	COMPONENT PUP3;	CHALIN: 1, W;	COMPONENT CITY CHAIN	T V. DBOTTE A COME	COMPONENT PREZ	CHAIN: K. Y:	PROTEASOME	COMPONENT CS; CHAIN:	L, Z; PROTEASOME	COMPONENT PRE4;	CLICALIVA, 184, 1.9
SEQ FOLD score	61.07																				,		-			-								
PMF score			0.25																								•							
Verify score			0.34	٠																•											<u> </u>		,	
Psi Blast	6.8e-31		3.4c-48		•														-															
END AA	139		295																									-					:	
START AA	21		69																				-											
CHAIN			А											_		_																		
PDB U	2rhe		1g0u																															
SEQ ID	426		428																															

Table 5

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PDB annotation	PROTEINASE YSCE SUBUNIT MACROPAIN SUBUNIT PRE3, PROTEINASE YSCE SUBUNIT PROTEASOME, UBIQUITIN, DEGRADATION, PROTEASE, NTN-	MULTICATALYTIC PROTEINASE MULTICATALYTIC PROTEINASE, 20S PROTEINASE, 20S PROTEASOME, PROTEIN 2 DEGRADATION, ANTIGEN PROCESSING, HYDROLASE, PROTEASE	MULTICATALYTIC PROTEINASE MULTICATALYTIC PROTEINASE, 20S PROTEINASP, PROTEIN 2 DEGRADATION, ANTIGEN PROCESSING, HYDROLASE, PROTEASE	MULTICATALYTIC PROTEINASE MULTICATALYTIC PROTEINASE, 20S PROTEINASE, 20S PROTEASOME, PROTEIN 2 (I) DEGRADATION, ANTIGEN (I) PROCESSING, HYDROLASE	MULTICATALYTIC PROTEINASE MULTICATALYTIC PROTEINASE, 208 PROTEASOME, PROTEIN 2 [1] DEGRADATION, ANTIGEN [1] PROCESSING, HYDROLASE [1]
Compound	PROTEASOME COMPONENT PRE3; CHAIN: N, 2;	20S PROTEASOME; CHAIN: A, B, C, D, B, F, G, H, I, J, K, L, M, N, O, P, Q,	20S PROTBASOME; CHAIN: A, B, C, D, E, F, G, H, I, J, K, L, M, N, O, P, Q,	20S PROTEASOME; CHAIN: A, B, C, D, E, F, G, H, I, I, K, L, M, N, O, P, Q,	20S PROTEASOME; CHAIN: A, B, C, D, E, F, G, H, I, I, K, L, M, N, O, P, Q,
SEQ FOLD score		76.38		·	66.44
PMF score		!	0.55	0.58	·
Verify			0.52	0.31	
Psi Blast		1.7e-47	1.7e-47	1.7e-48	1.7e-48
END AA		306	292	295	300
START		19	8	99	<i>L</i> 9
CHAIN		U	U .	四	E
PDB CD		1гур	Ігур	lryp	lryp
SEQ ID	·	428	428	428	428

Table 5

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PDB annotation	PROTEASE	MULTICATALYTIC PROTEINASE MULTICATALYTIC PROTEINASE, 20S	PROTEASOME, PROTEIN 2	PROCESSING, HYDROLASE, PROTEASE	MULTICATALYTIC PROTEINASE	MULTICATALYTIC PROTEINASE, 20S	PROTEASOME, PROTEIN 2 DEGRADATION ANTIGEN	PROCESSING, HYDROLASE, PROTEASE	MULTICATALYTIC PROTFINASE	MULTICATALYTIC	PROTEINASE, 20S	PROTEASOME, PROTEIN 2 DEGRADATION, ANTIGEN	PROCESSING, HYDROLASE, PROTEASE	YTIC	MULTICATALYTIC	PROTEINASE, 208	PROTEASOME, PROTEIN 2 U	PROCESSING, HYDROLASE	PROTEASE	PROTEINASE	YTIC		PROTEASOME, PROTEIN 2 FL DEGRADATION, ANTIGEN FL	J. C. C. C. C. C. C. C. C. C. C. C. C. C.
Сотроипа		20S PROTEASOME; CHAIN: A, B, C, D, E, F, G, H, I, J, K, L, M, N, O, P, Q,		-	20S PROTEASOME; CHAIN: A, B, C, D, E, F, G,	H, I, J, K, L, M, N, O, P, Q,			20S PROTEASOME;	H, I, J, K, L, M, N, O, P, Q,			-	20S PROTEASOME;	CHAIN: A, B, C, D, E, F, G, H, I, J, K, L, M, N, O, P, Q,					20S PROTEASOME;	H, I, I, K, L, M, N, O, P, Q,			
SEQ FOLD score					117.29									201.96					·					
PMF		1.00							1.00											00:				
Verify score		0.56							0.63						·					0.76				
Psi Blast		3.46-42			3.4e-42				8.5e-46				-	3.3e-56						3.3e-56				
END		273			328			,	295					305					-	309			· :	
START		95			95		·		95	-				95		_				S			<u></u>	
CHAIN		·			— ·			. 	ы					ı			1			<u>.</u>				
EOS II		lryp			lryp				lryp					lryp				,		1ryp				
SEQ ID	5	428			428				428					428						428				

Fable 5

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PDB annotation	PROCESSING, HYDROLASE, PROTEASE		COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC	FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX	(ZINC FINGERDINA), ZINC FINGER, DNA-BINDING PROTEIN				<i>/</i>				lano lano	- tt				Et.						GENE REGULATION POZ
Compound			QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX	OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C:	QGSR ZINC FINGER PEPTIDE; CHAIN: A;	DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B,	TRANSCRIPTION	REGULATION YEAST	ADRI (RESIDUES 102 - 130)	1ARD 3 (AMINO	TERMINAL ZINC FINGER	STRUCTURES) 1ARD 4	(ADRIB) 1ARD 5	DNA-BINDING PROTEIN	HUMAN ENHANCER-	MITTANT WITH CYS 11	1BBO 3 REPLACED BY	ABU (CIIABU) (NMR, 60	STRUCTURES) 1BBO 4	DNA-BINDING PROTEIN	HUMAN ENHANCEK- RINDING PROTRIN MRP-1	MUTANT WITH CYS 11	1BBO 3 REPLACED BY	ABU (C11ABU) (NIMR, 60	PROMYELOCYTIC
SEQ FOLD score																									
PMF			-0.01		-0.14		0.64	5						0.24						0.65					0.89
Verify			0.01		0.05		020	2						-0.41						-0.47				-	0.40
Psi			1e-28		1.2e-25		1 70 06	3						1.7e-11						1.3e-13					3.4e-19
END	900		120		573		303	070						520						520			_		170
START	WW .		471		498		907	6470						471						474					47
CHAIN			V		A										•							٠.			Ą
PDB			laih	·	lalh			DJB1						1bbo						1bbo					1buo
SEQ ID			430		430			084						430	-					430					430

Fable 5

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PDB annotation	DOMAIN; PROTEIN- PROTEIN INTERACTION POMAIN	DOMALY, TRANSCRIPTIONAL 2 REPRESSOR, ZINC-FINGER	PROTEIN, X-RAY CRYSTALLOGRAPHY, 3	PROTEIN STRUCTURE,	PROMYBLOCYTIC LEUKEMIA, GENE	REGULATION	COMPLEX (ZINC FINGER/DNA) ZINC FINGER,	PROTEIN-DNA	INTERACTION, PROTEIN	STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER,	PROTEIN-DNA	INTERACTION, PROTEIN	DESIGN, 2 CRISIAL	ZIRUCIUKE, CUMPLEA (ZINC FINGER/DINA)	COMPLEX (ZINC FINGER	PROTEIN-DNA	INTERACTION, PROTEIN	DESIGN, 2 CRYSTAL	STRUCTURE, COMPLEX	COMPLEX (ZINC	FINGER/DNA) ZINC FINGER,	PROTEIN-DINA INTERACTION PROTEIN	 STRUCTURE, COMPLEX (ZINC FINGER/DNA)	
Compound	LEUKEMIA ZINC FINGER PROTEIN PLZF; CHAIN: A;					:	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;	-			DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;				DNA; CHAIN: A, B, D, E;	PROTEIN: CHAIN: C. F. G.		·.		DNA: CHAIN: A. B. D. E.	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;		
SEQ FOLD score																										
PMF score							-0.20					0.22					-0.14					0 80	70.0			
Verify							0.11					-0.06					0.07		· ·			10 9				
Psi Blast							1.4e-45					1e-47					6.8e-42					1 40-11	11-24-11			
END							467					120					573					707	<u> </u>		:	
START							364			-	•	470					497				٠,	460	6 6			
CHAIN							υ υ					C					ပ					0	5		٠	
PDB CI	1						lmey					1mey	-				1mey			-			Ішеу			
SEQ ID						•	430					430					430					000	430			

Table :

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PDB annotation	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	ZINC FINGER TRANSCRIPTION FACTOR SP1; ZINC FINGER, TRANSCRIPTION ACTIVATION, SP1	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION INITIATION ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	TRANSCRIPTION REGULATION TRANSCRIPTION REGULATION, ADRI, ZINC FINGER, NMR		OXIDOREDUCTASE PROGESTERONE 21- HYDROXYLASE, CYPIICS P450 1, MEMBRANE PROTEIN, PROGESTERONE II 21-HYDROXYLASE, BENZO(A) 2 PYRENE
Compound	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;	SPIF2; CHAIN: NULL;	YYI; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	ADR1; CHAIN: NULL;	ZINC FINGER DNA BINDING DOMAIN ZINC- FINGER (ZFY-SWAP) (NMR, 12 STRUCTURES) 7ZNF 3	CYTOCHROMB P450 2C5; CHAIN: A;
SEQ FOLD score						
PMF	0.75	0.17	-0.15	0.40	0.16	1.00
Verify	0.02	-0.10	0.08	-0.43	-0.62	0.77
Psi Blast	1.4e-13	3.4c-09	16-30	5.1e-17	1e-05	0
END	522	526	573	524	526	497
START	495	498	477	474	498	55
CHAIN	O		υ U			A
PDB	Imey	1sp2	lubd	2adr	7znf	14t6
SEQ ID	430	430	430	430	430	433

Table

			·				
PDB annotation	HYDROXYLASE, ESTRADIOL 2- HYDROXYLASE, P450, CYP2C5	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING FROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING
Compound		QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C:	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE
SEQ FOLD score							·
PMF		0.39	0.74	0.75	0.59	0.72	0.71
Verify		-0.10	90.0	-0.10	-0.24	-0.24	0.00
Psi Rlast		le-23	1.7e-26	5.1e-30	6.8e-26	1.7e-23	16-25
END		337	365	628	655	894	923
START		244	286	544	572	830	842
CHAIN		V	ď	∢	∢	A	4
PDB		lalh	lalh	lalh	lalh	lalh	laih
SEQ ID		435	435	435	435	435	435

Table 5

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PDB annotation	PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN	COMPLEX (ZINC FINGER/DNA) COMPLEX (ZINC FINGER/DNA), ZINC FINGER, DNA-BINDING PROTEIN			NC FINGER ROTEIN FAL MPLEX	COMPLEX (ZINC FINGER/DNA) ZINC FINGER II PROTEIN-DNA INTERACTION, PROTEIN FIL
Compound	BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	QGSR ZINC FINGER PEPTIDE; CHAIN: A; DUPLEX OLIGONUCLEOTIDE BINDING SITE; CHAIN: B, C;	DNA-BINDING PROTEIN HUMAN ENHANCER- BINDING PROTEIN MBP-1 MUTANT WITH CYS 11 1BBO 3 REPLACED BY ABU (C11ABU) (NMR, 60 STRUCTURES) 1BBO 4	DNA-BINDING PROTEIN HUMAN ENHANCER- BINDING PROTEIN MBP-1 MUTANT WITH CYS 11 1BBO 3 REPLACED BY ABU (C11ABU) (NMR, 60 STRUCTURES) 1BBO 4	DNA; CHAIN; A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN; C, F, G;	DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN; CHAIN: C, F, G;
SEQ FOLD score							
PMF		0.04	0.36	0.46	0.45	0.59	0.99
Verify		-0.17	0.04	-0.40	-0.42	-0.03	-0.04
Psi		3.4e-24	le-26	8.5e-11	6.6e-14	3.46-42	5.1e-45
END		1030	1058	594	296	337	365
START	000	. 828	978	546	548	243	281
CHAIN	3	4	4			U	U
PDB	3	lalh	lalh	1bbo	1bbo	Imey	lmey
SEQ ID	Ž	435	435	435	435	435	435

			-			г ——		H=4	· IEme well-		Page 100	g	to Mark	mile bester.		4 .
PDB annotation	DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA	INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER, PROTEIN-DNA	INTERACTION, PROTEIN DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX CZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER,	PROTEIN-DNA INTERACTION, PROTEIN	DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX (ZINC FINGER/DNA)	COMPLEX (ZINC FINGER/DNA) ZINC FINGER	PROTEIN-DNA INTERACTION, PROTEIN	DESIGN, 2 CRYSTAL STRUCTURE, COMPLEX CTING HINGRE (IN A)	COMPLEX (ZINC FINGER, FINGER, ZINC FINGER, Z	PROTEIN-DNA INTERACTION. PROTEIN	DESIGN, 2 CRYSTAL FINAL STRUCTURE, COMPLEX		COMPLETO (ZELOC
Compound		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTFIN: CHAIN: C. F. G:		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER PROTEIN: CHAIN: C. F. G:		DNA; CHAIN: A, B, D, E; CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;		DNA; CHAIN: A, B, D, B; CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;		DNA; CHAIN: A, B, D, B;	PROTEIN; CHAIN: C, F, G;		TANK CHIAMI A D D D	DNA; CHAIN: A, B, D, E,
SEQ FOLD score				·												
PMF		0.42		0.39	·	0.83			0.39			0.03			,	0.86
Verify		-0.40		-0.23		-0.40			-0.15			-0.19				0.03
Psi	·	1.7e-40		1.7e-48		6.8e-35			1.2e-43			5.1e-42				3.4e-45
END		965		628		894			925			1030				1058
START		511		543		828	·		841			927				977
CHAIN		ပ		ပ		၁			၁			ပ				ပ
PDB		lmey		lmey		Imey			Imey			1mey				1mey
SEQ ID		435		435		435			435			435				435

Cable 5

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PDB annotation	FINGER/DNA) ZINC FINGER,	PROTEIN-DNA	INTERFECTION, FNOTEEN	CENTION & CANADI HY	(ZINC FINGER/DNA)	COMPLEX (ZINC	FINGER/DNA) ZINC FINGER,	PROTEIN-DNA	INTERACTION, PROTEIN	DESIGN, 2 CRYSTAL	STRUCTURE, COMPLEX	CONTRACTOR DAY (CONTRACTOR)	FINGER/DNA) ZINC FINGER.	PROTFIN-DNA	INTERACTION, PROTEIN	DESIGN 2 CRYSTAL	STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC	FINGER/DNA) ZINC FINGER,	PROTEIN-DNA	LEIN		STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (ZINC WINGER (DIA) ZINC FINGER	PPOTENTIAL DIA	INTERACTION, PROTEIN	DESIGN, 2 CRYSTAL	STRUCTURE, COMPLEX	(ZINC FINGER/DNA)	COMPLEX (TRANSCRIPTION	REGULATION/DIA) IFMA		TRANSCRIPTION FACTOR,
Compound	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;		-		DNA: CHAIN: A B D. B.	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;				7 4 4 7 4 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	DNA; CHAIN: A, B, D, E;	DECTRONS CHAIN OF A G.	rnolem, cirit.				DNA; CHAIN: A, B, D, E;	CONSENSUS ZINC FINGER	PROTEIN; CHAIN: C, F, G;	-				DNA; CHAIN: A, B, D, E;	CONSENSOS ZINC FINGER	FROIDING CITAIN: CALLS				TRANSCRIPTION FACTOR	MA; CHAIN: A; 5S RNA	deve, curan, e, .,	
SEQ FOLD				٠			·																												
PIMF						20.0	C7:0						-0.19				-		100	<u> </u>						0.94						0.45			
Verify	2025					0 50	°C')-						0.27						25 0-	}						0.44						-0.16			
Psi	Diast					10	1e-10			•			1.4e-08						8 50-11	1						1.2e-11						3.40-19	_		-
END	5					900	303						483						809	9	_					1002						628			
START	AA				•	62.0	6/7						456						507	750						976						544			
CHAIN	3					ļ	<u>۔</u> ق						G						C	ے ا						O	•					 	·		
PDB	1						lmey						1mey							I mey						Imey						4]		
SEQ ID	S.						435						435						200	433						435		,				425	3		

PDB annotation	5S RNA 2 GENE, DNA BINDING PROTEIN, ZINC FINGER, COMPLEX 3 (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATIONDNA) COMPLEX (TRANSCRIPTION REGULATIONDNA), RNA POLYMERASE III, 2 TRANSCRIPTION INITIATION, ZINC FINGER PROTEIN —	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING- YANG 1; TRANSCRIPTION INITIATION, INTIATOR ELEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA- PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING- TO YANG 1; TRANSCRIPTION PORTION, INTIATOR PLEMENT, YY1, ZINC 2 FINGER PROTEIN, DNA-PROTEIN RECOGNITION, 3 COMPLEX (TRANSCRIPTION REGULATION/DNA)	REGULATION/DNA) YING-YANG I; TRANSCRIPTION CONTIATION, INTIATION INTIATION CONTIANO COMPLEX (TRANSCRIPTION 31) COMPLEX (TRANSCRIPTION)
Compound		TFIIIA; CHAIN: A, D; 5S RIBOSOMAL RNA GENE; CHAIN: B, C, E, F;	YY1; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YY1; CHAIN: C; ADENO-ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;	YY1; CHAIN: C; ADBNO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT DNA; CHAIN: A, B;
SEQ FOLD score					·
PMF		0.09	0.80	09:0	0.36
Verify		-0.24	-0.34	-0.45	-0.39
Psi Rlast		1e-29	5.1e-29	5.16-26	6.8e-32
END		657	365	296	628
START		512	246	484	515
CHAIN		∢	U	ပ	ပ
PDB		1116	Ddul	lubd	Iubd
SEQ ID		435	435	435	435

Fable :

				-\					-			Τ.		IR	PE	- "	. ,100				1	* EDI	- }		12	12
FUD annotation	REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATIONDINA) YING-YANG 1; TRANSCRIPTION	ELEMENT, YY1, ZINC 2	FINGER PROTEIN, DNA-	COMPLEX (TRANSCRIPTION	REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-	YANG 1; TRANSCRIPTION INITIATION, INITIATOR	ELEMENT, YY1, ZINC 2	FINGER PROTEIN, DNA-	COMPLEX (TRANSCRIPTION	REGULATION/DNA)	COMPLEX (TRANSCRIPTION REGULATION/DNA) YING-YANG 1; TRANSCRIPTION	INITIATION, INITIATOR	FINGER PROTEIN, DNA-	PROTEIN RECOGNITION, 3	REGULATION/DNA)	TRANSCRIPTION	REGULATION TO A ANSCRIPTION	REGULATION, ADRI, ZINC	FINGER, NMR	TRANSCRIPTION F	TRANSCRIPTION	REGULATION, ADRI, ZINCIU		
Compound		YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT	DNA; CHAIN: A, B;			-	YY1; CHAIN: C; ADENO- ASSOCIATED VIRUS P5	INITIATOR BLEMENT			- 1		YYI; CHAIN: C; ADENO- ASSOCIATED VIRUS P5 INITIATOR ELEMENT	DNA; CHAIN: A, B;				ADRI; CHAIN: NULL;				ADRI; CHAIN: NULL;			COMPLEXATEANSCRIPTIO	
SEQ FOLD		·																		·						
PMF	21016	0.36		•			-0.19						0.70					0.88				0.24			9.0	10.19
Verify	SCUIC	-0.23					90.0						-0.21					-0.25] 			-0.34			100	20:0-
Psi	DIEST	6.8e-32					6.8e-12						1.7e-26					3 40-17				3.4e-15			3	9.96-15
END	₩.	655					169						921					508	3	·		630				296
START	AA	551					611						837					546) 			572				545
CHAIN		υ		•			Ú						ပ													4
	e	lubd					lubd						lubd					4.00	7801			2adr				2drp
SEQ ID	NO:	435					435						435		·			,	435			435				435

			· · · · · · · · · · · · · · · · · · ·	 -T	PETM		
PDB annotation				COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE- FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA- BINDING PROTEIN/DNA)	COMPLEX (DNA-BINDING PROTEIN/DNA) FIVE- TO FINGER GLI; GLI, ZINC FINGER, COMPLEX (DNA-BINDING PROTEIN/DNA)	SIGNALING PROTEIN PHOTORECEPTOR, G PROTEIN-COUPLED RECEPTOR, MEMBRANE PROTEIN, 2 RETINAL PROTEIN, VISUAL PIGMENT	
Compound	N REGULATIONDNA) TRAMTRACK PROTEIN (TWO ZINC-FINGER PEPTIDE) COMPLEXED WITH 2DRP 3 DNA 2DRP 4	COMPLEX(TRANSCRIPTIO N REGULATION/DNA) TRAMTRACK PROTEIN (TWO ZINC-FINGER PEPTIDE) COMPLEXED WITH 2DRP 3 DNA 2DRP 4	COMPLEX(TRANSCRIPTIO N REGULATION/DNA) TRAMTRACK PROTEIN (TWO ZINC-FINGER PEPTIDE) COMPLEXED WITH 2DRP 3 DNA 2DRP 4	ZINC FINGER PROTEIN GLII; CHAIN: A; DNA; CHAIN: C, D;	ZINC FINGER PROTBIN GLII; CHAIN: A; DNA; CHAIN: C, D;	RHODOPSIN; CHAIN: A, B	GROWTH FACTOR ACIDIC FIBROBLAST GROWTH FACTOR (AFGF) MUTANT WITH CYS 47 1AFC 3
SEQ FOLD score							51.39
PMF		0.33	0.52	0.58	0.65	0.01	
Verify		-0.46	-0.30	-0.44	-0.26	-0.47	
Psi		2.6e-16	6.8e-08	8.5e-25	1.5e-29	1.4e-11	6.8e-33
END		628	627	394	657	300	174
START		567	570	250	516	83	38
CHAIN		4	4	V		æ	4
PDB		2drp	2drp	2gli	2gli	1f88	lafc
SEQ ID		435	435	435	435	874	437

· .	· · · · ·		Γ		T. T. WSI	EZULEE	
PDB annotation							GROWTH FACTOR FGF-2;
Compound	REPLACED BY ALA (C47A) COMPLEX WITH SUCROSE OCTASULFATE 1AFC 4	GROWTH FACTOR ACIDIC FIBROBLAST GROWTH FACTOR (AFGF) MUTANT WITH CYS 47 1AFC 3 REFLACED BY ALA (C47A) COMPLEX WITH SUCROSE OCTASULFATE 1AFC 4	GROWTH FACTOR ACIDIC FIBROBLAST GROWTH FACTOR (AFGF) MUTANT WITH CYS 47 1BAR 3 REPLACED BY ALA AND HIS 93 REPLACED BY GLY (C47A, H93G) 1BAR 4	GROWTH FACTOR ACIDIC FIBROBLAST GROWTH FACTOR (AFGF) MUTANT WITH CYS 47 1BAR 3 REPLACED BY ALA AND HIS 93 REPLACED BY GLY (C47A, H93G) 1BAR 4	GROWTH FACTOR BASIC FIBROBLAST GROWTH FACTOR MUTANT WITH CYS 69 REPLACED 1BFG 3 BY SER AND CYS 87 REPLACED BY SER (C69S,C87S) 1BFG 4	GROWTH FACTOR BASIC FIBROBLAST GROWTH FACTOR MUTANT WITH CYS 69 REPLACED 1BFG 3 BY SER AND CYS 87 REPLACED BY SER (C69S,C87S) 1BFG 4	BASIC FIBROBLAST
SEQ FOLD score			55.74		56.94		67.74
PMF score		0.64		0.76		0.46	
Verify score		0.38		0.56		0.61	
Psi Blast		6.8e-33	1.7e-33	1.7e-33	3.4e-36	3.4e-36	1.7e-39
END AA	•	173	174	173	164	172	164
START AA	-	45	82	33	38	4	=
CHAIN ID		4	æ	æ			
PDB ID		1afc	Ibar	1bar	1bfg	1bfg	1bla
SEQ ID NO:	·	437	437	437	437	437	437

Table 5

			_		_	_	_		_							
PDB annotation		GROWTH FACTOR	GROWTH FACTOR FGF-2;	GROWTH FACTOR	TANK TO TO TO THE OWNER OF THE OWNER OWN	HOKMONEGROWIN	FACTOR BETA-TREFOIL	HORMONE/GROWTH	FACTOR BETA-TREFOIL	HORMONE/GROWTH	FACTOR BETA-TREFOIL,	HORMONE/GROWTH	FACTOR	GROWTH FACTOR AFG;	2AFG 6	
Compound		GROWTH FACTOR; CHAIN: NUIL:	BASIC FIBROBLAST	GROWTH FACTOR;	THE ROLL OF CHANGE	FIBROBLAST GROWTH	FACTOR 7; CHAIN: A, B;	FIBROBLAST GROWTH	PACTOR 7; CHAIN: A, B;	FIBROBLAST GROWTH	FACTOR 7/1 CHIMERA;	CHAIN: A;		ACIDIC FIBROBLAST	GROWTH FACTOR; 2AFG 4	CHAIN: A, B, C, D; 2AFG 5
CTO4 DES	score					•										
PMF	score		0.03		72.0	9/0		0.25		0.75				99.0		
Verify	score		0.11		9,	0.40		0.29		0.62				0.49	-	
Psi	Blast		1.7e-39 0.11	•	,	3.46-41 0.40		1.5e-37		5.1e-37		•		1.2e-33		
END	Ψ¥		172		Ş	1/2		172		173				173		
CHAIN START	¥		16		ļ	35		39		39				45		
CHAIN	A				•	∢		В		A		-		V.	,	
PDB	A		1bla			Idqk		lqqk		lggl				2afg		
SEQ ID PDB	ÖN		437		50,	43/		437		437				437		

Table 6

SEQ.ID NO:	Position of Signal Peptide	Maximum score	Mean score	
1	24	0.978	0.760	
2	32	0.995	0.681	
3	37	0.979	0.718	
4	18	0.925	0.822	
5	28	0.939	0.749	
6	41	0.989	0.690	
7	26	0.960	0.674	
8	16	0.973	0.925	
9	24	0.978	0.760	
10	18	0.887	0.579	
• 11	42	0.977	0.587	
12	21	0.966	0.848	
13	25	0.993	0.954	
14	28	0.909	0.664	
16	23	0.913	0.597	
17	42	0.978	0.689	
18	21	0.930	0.662	
19	45	0.985	0.714	
20	37	0.983	0.855	
21	31	0.947	0.833	
22	20	0.979	0.773	
24	30	0.924	0.720	
25	26	0.974		
26	28		0.824	
28	16	0.982	0.649	
29	27	0.912	0.705	
30	22	0.957	0.652	
	 	0.968	0.844	
31	23	0.952	0.812	
32	18	0.932	0.884	
33	29	0.991	0.729	
34	26	0.939	0.709	
35	29	0.961	0.842	
36	16	0.951	0.777	
37	27	0.983	0.898	
38	17	0.991	0.955	
39	33	0.977	0.822	
40	17	0.989	0.969	
41	30	0.936	0.679	
42	24	0.993	0.810	
44	22	0.990	0.921	
54.	18	0.925	0.822	
56	18	0.981	0.951	
60	28	0.939	0.749	
62	33	0.979	0.757	
70	41	0.989	0.690	
79	26	0.960	0.674	
83	18	0.979	0.963	
84	22	0.967	0.792	
87	25	0.980	0.867	
97	16	0.973	0.925	
98	24	0.978	0.760	
99	17	0.978	0.925	

Table 6

SEQ.ID NO: Position of Signal Peptide		Maximum score	Mean score
113	18	0.887	0.579
115	18	0.952	0.670
120	42	0.977	0.587
137	21	0.966	0.848
140	25	0.993	0.954
153	28	0.909	0.664
156	18	0.954	0.747
174	23	0.913	0.597
175	20	0.986	0.936
178	42	0.978	0.689
180	32	0.929	0.583
184	21	0.979	0.941
192	21	0.930	0.662
200	45	0.985	0.714
212	37	0.983	0.855
225	24	0.992	0.882
228	20		
237	17	0.979	0.911
251	13	0.982	0.964
		0.918	0.692
252	13	0.918	0.692
256	20	0.912	0.693
257	20	0.912	0.693
260	26	0.974	. 0.824
262	18	0.965	0.833
267	25	0.956	0.765
288	16	0.912	0.705
289	18	0.896	0.634
290	19	0.966	0.897
294	18	0.991	0.973
295	20	0.906	0.580
299	27	0.957	0.652
307	19	0.983	0.871
310	22	0.968	0.844
320	23	0.952	0.812
324	27	0.982	0.911
327	18	0.983	0.941
328	18	0.932	0.884
332	27	0.990	0.923
335	45	0.983	0.793
336	45	0.983	0.793
346	29	0.991	0.729
354	22	0.978	0.877
363	26	0.939	0.709
364	22	0.966	0.843
375	29	0.961	0.842
379	16	0.951	0.777
401	44	0.975	0.876
407	33	0.977	0.822
417	17	0.989	0.969
418	23	0.974	0.799
422	18	0.981	0.952
426	21	0.982	0.912

Table 6

SEQ.ID NO:	Position of Signal Peptide	Maximum score	Mean score
428	30	0.936	0.679
429	43	0.978	0.712
433	28	0.993	0.948
434	43	0.930	0.624
437	24	0.993	0.810
438	16	0.978	0.939

Table 7

SEQ ID NO:	Chromsomal location
3	2q11.2
4	20pter-p12.3
5	5q31
6	19p12
7	19p12
8	5
11	12p13-p12
12	12p13-p12 p11.2-12.3
13	19p
14	6p12.1-21.1
15	19p13.1
17	16q12-q13
19	15
20	15 15
22	Xq13.1
23	12
25	11p15.5
26	20
27	22
28	12q23-24.1
29	20
30	13
31	12
33	15
36	4q28
37	14q24.3
38	10
39	20
41	17q12-q21
42	14
44	1q24.1-25.2
45	2
47	
48	3q21-q25 9
49	14
50	6q14.1-15
51	19
. 52	11
53	20
54	16
55	14
56	3
57	19
58	19 7p15.1-p13
59	19
61	19 2
62	19
63	16
66	15
70	1p31.1-33
71	9
72	16

Table 7

SEQ ID NO:	Chromsomal location
74	5q31-q33
75	3p21.1-q13.13
76	3p21.1-q13.13 2
77	2
78	21q22.1
79	Xp11.22-p11.21
80	2
81	19
82	20
83	19p13.3
84	19
85	3
86	8
87	1p13
88	16
89	18q21.1-q22
90	11q13.1-q13.3
91	18p11.23-p11.21
92	17
93	10
94	3
95	x
96	6q14.2-16.1
97	1q21.2-22
98	1q21.2-22
99	66_
102	8q22-q23
103	10p11.2
104	17
105	17
106	2
107	1
108	16
109	17q21.3-q22
110	11q
111	3p21.1-q13.13
112	16
113	5
114	9
115	3p13-q26.1
116	3p13-q26.1 5 7q31
117	7q31
118	14
119	14
120	19
121	19
121 122	6q27
123	6q27 14
124	1g21-g22
125	1q21-q22 6
126	17q25
127	15
	

Table 7

SEQ ID NO:	Chromsomal location
129	14q31
130	1p36.1
131	11
132	20
133	20p11.23-p11.21
134	1p32
135	2031
136	2q31 X
138	12p13
139	9
140	p34.1-34.3
140	19q12
142	
	15q26
143	22q11.21
144	17q12
145	4p16.3
146	22
147	16p11.2
148	18q12
150	4
151	7p12-q11.21
152	14
153	14q32.33
155	1p34
156	16p13.3
157	12p13.3
158	5
159	8
160	19
161	4
162	1
163	11q23
164	3
165	12q22
168	19
170	1
171	18q12
173	7
174	13
175	2p23.3-q32.3
176	2p23.3-q32.3 16
178	10
179	1q21-q25
180	19p13.3
181	1
184	1p35.1-36.23
185	1953.1-30.25
186	18
187	
188	3p13-q26.1
100	3
189	17
190	6

Table 7

SEQ ID NO:	Chromsomal location
193	11p15.5
194	14q32
195	12
196	10q24
198	1p36.1
199	5q22
200	11
201	2q31
202	17
206	Xp11.23
207	Ap11.25
208	9q34 19
208	20
210	11q23
211	16p12
212	19q13.1
213	7p15
214	15
215	1p36.21-36.33
216	11
217	22q11.2
- 218	15
219	19q13.4
222	19
223	1q25.2
226	1
227	1p36.11-36.23
228	1p36.3-p36.13
230	17
231	7q33-q34
232	3
233	9
234	10
235	17
236	4
237	19q13.4
238	4q25
239	2
240	7
241	12
243	6p21.3
244	3p13-q26.1
245	3913-420.1
245	1 1 2 4 1
246	1p34.1
247	3q23
	3p21.3
249	20
250	20
251	18q12-q21
252	18q12-q21
253	14
254	1p35.3-p35.1

Table 7

SEQ ID NO:	Chromsomal location
256	6q25-q26
257	6q25-q26
258	1q21-q23
259	16p13.2-16p13.11
260	14q21.1-q24.1
261	2p23.3-q32.3
262	12
263	19
264	4q28
265	
266	2 2
	1
267	1q21-q23
268	20p12.3-p13
269	4
270	6
271	2p23.3-q14.3
272	18q21
273	18q21
274	14q22
275	6p21.3
276 .	5
280	8
281	4q22-q24
282	2
283	7q22-q31.1
284	11
285	11q12.3
286	10
287	19
290	17
291	4q22 ·
292	1p36.11-36.23
293	19
294	22
296	3
297	4p16
298	6
299	8q13
300	20
301	15
302	22q11.2-q22
303	15
303	6
306	6
306	
307	9p24.2
	2p23.3-q24.3
309	14
310	6
311	2
312	4
313	19pter-19p13.3 3
314]3

Table 7

SEQ ID NO:	Chromsomal location
316	11p12-14.2
317	19
318	17
319	17
320	5q14
323	4
324	3p
325	6p21.1-21.31
326	17p11.2
327	9
328	5q23
329	2
330	3
331	1p21.1-22.1
332	9 7
333	
334	11q13
337	14
338	7q35-q36
339	13
340	6q11.1-22.33
341	11q12-q13.1
343	10
344	16
345	16
346	11q22
347	19
348	15q24-q26
350	Xp11.21-11.22
354	16
355	19
356	11
358	Xp11.23
359	4
360	8
362	4
363	11
364	11g13
365	7q31
366	22q13.31-13.32
367	5
370	19
371	7q31.1-7q31.33
372	2q37.3
373	3
374	16
375	19q13.4
376	18q12
377	18q12
377	8
	11q13
380	6
381	0

Table 7

SEQ ID NO:	Chromsomal location
385	4q28
386	15
387	10
388	17
389	11p15.4
390	6p21.3
391	22q13
392	3
393	
394	15
395	1
396	6p21.2-p21.3
397	15
399	7q31
400	14
402	Xq28
403	10
404	16
406	16
408	11
412	20q12-13.1
413	15
414	17
415	4
416	12q
419	21q22.1
420	16p11.2 6
422	6
424	21
426	14
428	14
429	1q22-q23
430	11q13
431	3
432	2
433	19q13.1
434	20q13.1
435	18q23
436	11q24
437	10
438	4q21-q25

Table 8

SEQ ID NO: of Full-length Nucleotide Sequence	SEQ ID NO: of Full-length Nucleotide Sequence	SEQ ID NO: in Priority Application USSN 09/774,528	
52	52	54	
53	53	55	
54	54	56	
55	55	57	
56	56	58	
57	57	59	
58	58	60	
59	59	61	
60	60	62	
61	61	63	
62	62	64	
63	63	65	
64	64	66	
65	65	67	
66	66	68	
67	67	69	
68	68	70	
69	69		
70		71	
	70	12	
71	71	73	
72	72	74	
73	73	75	
74	74	76	
75	75	77	
76	76	78	
77	77	79	
78	78	80	
79	79	81	
80	80	82	
81	81	83	
82	82	. 84	
83	83	85	
84	84	86	
85	85	87	
86	86	88	
87	87	89	
88	88	.90	
89	89	91	
90	90	92	
91	91	93	
92	92	94	
93	93	95	
94	94	96	
95	95	97	
96	96	98	
97	97 .	99	
98	98	100	
99	99		
100		101	
	100		
101	101	103	
102	102	104	
103	103	105	

Table 8

SEQ ID NO: of Full-length Nucleotide Sequence	SEQ ID NO: of Full-length Nucleotide Sequence	SEQ ID NO: in Priority Application USSN 09/774,528	
104	104	106	
105	105	107	
106	106	108	
107	107	109	
108	108	110	
109	109	111	
110	110	112	
111	111	113	
112	112	114	
113	113	115	
114	114	116	
115	115	117	
116	116	118	
117	. 117	119	
118	118	120	
119	119	121	
120	120	122	
121	121	123	
122	122	124	
123	123	125	
124	124	126	
125	125	127	
126	126	128	
127	127	129	
128	128	130	
129	129	131	
130	130	132	
131	131	133	
132	132	134	
133	133	135	
134	134	136	
135	135	137	
136	136	138	
137	137	139	
138	138	140	
139	139	141	
140	140	142	
141	141	143	
142	142	144	
143	143	145	
144	144	146	
145	145	147	
146	146	148	
147	147	148	
148	148	150	
149	149	151	
150	150	151	
151	151	152	
151	152	153	
152			
153	153 154	155 156	

Table 8

SEQ ID NO: of Full-length	SEQ ID NO: of Full-length	SEQ ID NO: in Priority Application	
Nucleotide Sequence	Nucleotide Sequence	USSN 09/774,528	
156	156	158	
157	157	159	
158	158	160	
159	159	161	
160	160	162	
161	161	163	
162	162	164	
163	163	165	
164	164	166	
165	165	167	
166	166	168	
167	167	169	
168	168	170	
169	169	171	
170	170	172	
171	171	173	
172	172	174	
173	173	175	
174	174	176	
175	175	177	
176	. 176	178	
177	177	179	
178	178	180	
179	179	181	
180	180	182	
181	181	183	
182	182	184	
183	183	185	
184	184	186	
185	185	187	
186	186	188	
. 187	187	189	
188	188	190	
189	189	191	
190	190	192	
191	191	193	
192	. 192	194	
193	193	195	
194	194	196	
195	195	197	
196	196	198	
197	197	199	
198	198	200	
199	199	201	
200	200	202	
201	201	203	
202	202	204	
203	203	205	
204	204	206	
205	205	207	
206	206	208	
207	207	209	

Table 8

SEQ ID NO: of Full-length Nucleotide Sequence	ull-length SEQ ID NO: of Full-length quence Nucleotide Sequence	SEQ ID NO: in Priority Application USSN 09/774,528	
208	208	210	
209	209	211	
210	210	212	
211	211	213	
212	212	214	
213	213	215	
214	214	216	
215	215	217	
216	216	218	
217	217	219	
218	218	220	
219	219	221	
220	220	222	
221	221	223	
222	222	224	
223	223	225	
224	224	225	
225	225	226	
226	226		
227	227	228	
228		229	
	228	230	
229	229	231	
230	230	232	
231	231	233	
232	232	234	
233	233	235	
234	234	236	
235	235	237	
236	236	238	
237	237	239	
238	238	240	
239	239	241	
240	240	242	
241	241	243	
242	242	244	
243	243	245	
. 244	244	246	
245	245	247	
246	246	248	
247	247	249	
248	248	250	
249	249	251	
250	250	252	
251	251	253	
252	252	254	
253	253	255	
254	254	256	
255	255	257	
256	256	258	
257	257	258	
258	258	259	

Table 8

SEQ ID NO: of Full-length Nucleotide Sequence	SEQ ID NO: of Full-length Nucleotide Sequence	SEQ ID NO: in Priority Application USSN 09/774,528
260	260	262
261	261	263
262	262	264
263	263	265
264	264	266
265	265	267
266	266	268
267	267	269
268	268	270
269	269	271
270	270	272
271	271	273
272	272	274
273	273	275
274	274	276
275	275	277
276	276	278
277	277	279
278	278	200
279	279	281
280	280	282
281	281	283
282	282	284
283	283	285
284	284	286
285	285	287
286	286	288
287	287	289
288	288	290
289	289	291
290	290	292
291	291	293
292	292	294
293	293	295
294	294	296
295	295	297
296	296	298
297	297	299
298	298	300
299	299	301
300	300	302
301	301	303
302	302	304
303	303	305
304	304	306
305	305	307
306	306	308
307	307	309
308	308	310
309	309	311
310	310	312
311	311	313
	<u> </u>	

Table 8

SEQ ID NO: of Full-length	SEQ ID NO: of Full-length	SEQ ID NO: in Priority Application	
Nucleotide Sequence	Nucleotide Sequence	USSN 09/774,528	
312	312	314	
313	313	315	
314	314	316	
315	315	317	
316	316	318	
317	317	319	
318	318	320	
319	319	321	
320	320	322	
321	321	323	
322	322	324	
323	323	325	
324	324	326	
325	325	327	
326	326	328	
327	327	329	
328	328	330	
329	329	331	
330	330	332	
331	331	333	
332	332	334	
333	333	335	
334	334	336	
335	335	337	
336	336	338	
337	337	339	
338	338	340	
339	339	341	
340	340	342	
341	341	343	
342	342	344	
343	343	345	
344	344	346	
345	345	347	
346	346	348	
347	347	349	
348	348	350	
349	349	351	
350	350	352	
351	351	353	
352	352	354	
353	353	355	
354	354	356	
355 .	355	357	
356	356	358	
357	357	360	
358	358	361	
359	359	362	
360	360	363	
361	361	364	
362	362	365	
363	363	366	
	1. 303	1 300	

Table 8

SEQ ID NO: of Full-length	SEQ ID NO: of Full-length	SEQ ID NO: in Priority Application	
Nucleotide Sequence	Nucleotide Sequence	USSN 09/774,528	
364	364	367	
365	365	368	
366	366	369	
367	367	370	
368	368	371	
369	369	372	
370	370	373	
371	371	374	
372	372	375	
373	373	376	
374	374	377	
375	375	378	
376	376	379	
377	377	380	
378	378	381	
379	379	382	
380	380	383	
381	381	384	
382	382	385	
383	383	386	
384	384	387	
385	385	388	
386	386	389	
387	387		
388	388	390	
389	389	391	
390	390	392	
390		393	
391	391	394	
	392	395	
393	393	396	
394	394	397	
395	395	398	
396	396	399	
397	397	400	
398	398	401	
399	399	402	
400	400	403	
401	401	404	
402	402	405	
403	403	406	
404	404	407	
405	405	408	
406	406	409	
407	407	410	
408	408	411	
409	409	412	
410	410	413	
411	411	414	
412	412	415	
413	413	416	
414	414	417	
415	415	418	

Table 8

SEQ ID NO: of Full-length	SEQ ID NO: of Full-length	SEQ ID NO: in Priority Application	
Nucleotide Sequence	Nucleotide Sequence	USSN 09/774,528	
416	416	419	
417	417	420	
418	418	421	
419	419	422	
420	420	423	
421	421	424	
422	422	425	
423	423	426	
424	424	427	
425	425	428	
426	426	429	
427	427	430	
428	428	431	
429	429	432	
430	430	433	
431	431	434	
432	432	435	
433	433	436	
434	434	437	
435	435	438	
436	436	439	
437	437	440	
438	438	441	

WHAT IS CLAIMED IS:

- 1. An isolated polynucleotide comprising a nucleotide sequence selected from the group consisting of SEQ ID NO: 1-438, a mature protein coding portion of SEQ ID NO: 1-438, an active domain coding portion of SEQ ID NO: 1-438, and complementary sequences thereof.
- 2. An isolated polynucleotide encoding a polypeptide with biological activity, wherein said polynucleotide hybridizes to the polynucleotide of claim 1 under stringent hybridization conditions.
- 3. An isolated polynucleotide encoding a polypeptide with biological activity, wherein said polynucleotide has greater than about 90% sequence identity with the polynucleotide of claim 1.
- 4. The polynucleotide of claim 1 wherein said polynucleotide is DNA.
- 5. An isolated polynucleotide of claim 1 wherein said polynucleotide comprises the complementary sequences.
- 6. A vector comprising the polynucleotide of claim 1.
- 7. An expression vector comprising the polynucleotide of claim 1.
- 8. A host cell genetically engineered to comprise the polynucleotide of claim 1.
- 9. A host cell genetically engineered to comprise the polynucleotide of claim 1 operatively associated with a regulatory sequence that modulates expression of the polynucleotide in the host cell.
- 10. An isolated polypeptide, wherein the polypeptide is selected from the group consisting of:

(a) a polypeptide encoded by any one of the polynucleotides of claim 1; and

- (b) a polypeptide encoded by a polynucleotide hybridizing under stringent conditions with any one of SEQ ID NO: 1-438.
- 11. A composition comprising the polypeptide of claim 10 and a carrier.
- 12. An antibody directed against the polypeptide of claim 10.
- 13. A method for detecting the polynucleotide of claim 1 in a sample, comprising:
- a) contacting the sample with a compound that binds to and forms a complex with the polynucleotide of claim 1 for a period sufficient to form the complex;
 and
- b) detecting the complex, so that if a complex is detected, the polynucleotide of claim 1 is detected.
- 14. A method for detecting the polynucleotide of claim 1 in a sample, comprising:
- a) contacting the sample under stringent hybridization conditions with nucleic acid primers that anneal to the polynucleotide of claim 1 under such conditions;
- b) amplifying a product comprising at least a portion of the polynucleotide of claim 1; and
- c) detecting said product and thereby the polynucleotide of claim 1 in the sample.
- 15. The method of claim 14, wherein the polynucleotide is an RNA molecule and the method further comprises reverse transcribing an annealed RNA molecule into a cDNA polynucleotide.
- 16. A method for detecting the polypeptide of claim 10 in a sample, comprising:

 a) contacting the sample with a compound that binds to and forms a complex with the polypeptide under conditions and for a period sufficient to form the complex; and

- b) detecting formation of the complex, so that if a complex formation is detected, the polypeptide of claim 10 is detected.
- 17. A method for identifying a compound that binds to the polypeptide of claim 10, comprising:
- a) contacting the compound with the polypeptide of claim 10 under conditions sufficient to form a polypeptide/compound complex; and
- b) detecting the complex, so that if the polypeptide/compound complex is detected, a compound that binds to the polypeptide of claim 10 is identified.
- 18. A method for identifying a compound that binds to the polypeptide of claim 10, comprising:
- a) contacting the compound with the polypeptide of claim 10, in a cell, under conditions sufficient to form a polypeptide/compound complex, wherein the complex drives expression of a reporter gene sequence in the cell; and
- b) detecting the complex by detecting reporter gene sequence expression, so that if the polypeptide/compound complex is detected, a compound that binds to the polypeptide of claim 10 is identified.
- 19. A method of producing the polypeptide of claim 10, comprising,
- a) culturing a host cell comprising a polynucleotide sequence selected from SEQ ID NO: 1-438, a mature protein coding portion of SEQ ID NO: 1-438, an active domain coding portion of SEQ ID NO: 1-438, complementary sequences thereof and a polynucleotide sequence hybridizing under stringent conditions to SEQ ID NO: 1-438, under conditions sufficient to express the polypeptide in said cell; and
 - b) isolating the polypeptide from the cell culture or cells of step (a).

20. An isolated polypeptide comprising an amino acid sequence selected from the group consisting of any one of the polypeptides encoded by SEQ ID NO: 1-438, the mature protein portion thereof, or the active domain thereof.

- 21. The polypeptide of claim 20 wherein the polypeptide is provided on a polypeptide array.
- 22. A collection of polynucleotides, wherein the collection comprising the sequence information of at least one of SEQ ID NO: 1-438.
- 23. The collection of claim 22, wherein the collection is provided on a nucleic acid array.
- 24. The collection of claim 23, wherein the array detects full-matches to any one of the polynucleotides in the collection.
- 25. The collection of claim 23, wherein the array detects mismatches to any one of the polynucleotides in the collection.
- 26. The collection of claim 22, wherein the collection is provided in a computerreadable format.
- 27. A method of treatment comprising administering to a mammalian subject in need thereof a therapeutic amount of a composition comprising a polypeptide of claim 10 or 20 and a pharmaceutically acceptable carrier.
- 28. A method of treatment comprising administering to a mammalian subject in need thereof a therapeutic amount of a composition comprising an antibody that specifically binds to a polypeptide of claim 10 or 20 and a pharmaceutically acceptable carrier.

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- (22) International Filing Date: 29 January 2002 (29.01.2002)
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English

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- (81) Designated States (national): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZM, ZW.
- (84) Designated States (regional): ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).

Published:

- with international search report
- with sequence listing part of description published separately in electronic form and available upon request from the International Bureau
- (88) Date of publication of the international search report: 5 August 2004

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.



International application No.

PCT/US02/01222

A. CLASSIFICATION OF SUBJECT MATTER IPC(7) : C12N 9/00, 1/20, 15/00; C12P 21/04; C07H 21/04 US CL : 435/183, 252.3, 320.1, 71.1; 536/23.2 According to International Patent Classification (IPC) or to both national classification and IPC				
B. FIELDS SEARCHED				
Minimum documentation searched (classification system followed U.S.: 435/183, 252.3, 320.1, 71.1; 536/23.2	Minimum documentation searched (classification system followed by classification symbols)			
Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched				
Electronic data base consulted during the international search (name of data base and, where practicable, search terms used) STN/EAST, Est database, Strembl database, PIR database				
C. DOCUMENTS CONSIDERED TO BE RELEVANT				
Category * Citation of document, with indication, where a	<u> </u>			
X EP 1 003 401 A2 (DUMAS et al) 06 Sepember 200	0 (06.09.2000). 1-9, 19 and 22-26			
Further documents are listed in the continuation of Box C.	See patent family annex.			
* Special categories of cited documents: "T" later document published after the international filling date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention				
"E" earlier application or patent published on or after the international filing date	"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone			
"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"Y" document of particular relevance; the claimed invention cannot be considered to one or more other such document is combined with one or more other such documents, such combination			
"O" document referring to an oral-disclosure, use, exhibition or other means being obvious to a person skilled in the art				
"P" document published prior to the international filing date but later than the "&" document member of the same patent family priority date claimed				
Date of the actual completion of the international search Date of mailing of the international search report				
04 December 2003 (04.12.2003) 18 APR 2004				
Name and mailing address of the ISA/US Mail Stop PCT, Attn: ISA/US Commissioner for Patents Authorized officer Ponnathapu Achutamurthy				
P.O. Box 1450 Alexandria, Virginia 22313-1450 Telephone No. 703-308-0196				
Facsimile No. (703)305-3230 Form PCT/ISA/210 (second sheet) (July 1998)				

International application No.

PCT/US02/01222

Box I Observations where certain claims were found unsearchable (Continuation of Item 1 of first sheet)			
This international report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:			
1. Claim Nos.: because they relate to subject matter not required to be searched by this Authority, namely:			
Claim Nos.: because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:			
3. Claim Nos.: because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).			
Box II Observations where unity of invention is lacking (Continuation of Item 2 of first sheet)			
This International Searching Authority found multiple inventions in this international application, as follows: Please See Continuation Sheet			
As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.			
2. As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.			
3. As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:			
. K-7			
4. No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.: 1-9, 19 and 22-26 (SEQ ID NO:231)			
Remark on Protest The additional search fees were accompanied by the applicant's protest.			
No protest accompanied the payment of additional search fees.			

PCT/	US02/	012	22

BOX II. OBSERVATIONS WHERE UNITY OF INVENTION IS LACKING

This application contains the following inventions or groups of inventions which are not so linked as to form a single general inventive concept under PCT Rule 13.1. In order for all inventions to be examined, the appropriate additional examination fees must be paid.

Group I, claim(s) 1-9,19 and 22-26, drawn to DNA of SEQ ID NO:1-438, vector comprising said DNA, host cell comprising said DNA and a method of producing polypeptides.

Group II, claim(s) 10-11 and 20-22, drawn to polypeptides encoded by the DNA of Group I.

Group III, claim(s) 12, drawn to antibody against the protein of Group II.

Group IV, claim(s) 13-15, drawn to a method of detecting the DNA of Group I.

Group V, claim(s) 16, drawn to a method of detecting the polypeptide of Group II.

Group VI, claim(s) 17-18, drawn to a method of identifying a compound that bind to the polypeptide of Group II.

Group VII, claim(s) 27 drawn to a method of treatment using the polypeptide of Group II.

Group VIII, claim(s) 28, drawn to a method of treatment using the antibody of Group III.

The inventions listed as Groups I-VIII do not relate to a single general inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, they lack the same or corresponding special technical features for the following reasons:

The DNA of SEQ ID NO:1-438 are different in structure and encode polypeptides having different structure and different function or substrate specificity. Therefore, in addition to electing one Group, applicants must further elect one DNA sequence or one polypeptide sequence encoded by SEQ ID NO:1-438.

The technical feature linking Groups I-VIII appears to be that they all relate to the DNA of SEQ ID NO:1-438. However, Dumas et al. teach a polypeptide encoded by a polymcleotide that is 99% identical to SEQ ID NO:231.

Therefore, the technical feature linking the inventions of Groups I-X does not constitute a special technical feature as defined by PCT Rule 13.2, as it does not define a contribution over the prior art.

Groups I-III do not share a technical feature because a DNA, a protein, and an antibody are different compounds, each with its own chemical structure and function, and they have different utilities. The DNA molecule of Group I is not limited in use to the production of polypeptide of Group II and can be used as a hybridization probe, and protein of Group II can be obtained by a materially different method such as by biochemical purification. The structure of an antibody of Group III is not predictable from the structure of the protein of Group II and an antibody can cross-react with various proteins.

The special technical feature of Group I is a DNA of SEQ ID NO:1-438, vector comprising said DNA, host cell comprising said DNA and a method of producing polypeptides.

The special technical feature of Group II is a polypeptide encoded by the DNA of Group I.

The special technical feature of Group III is an antibody against the protein of Group II.

The special technical feature of Group IV is a a method of detecting the DNA of Group I.

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The special technical feature of Group V is a a method of detecting the polypeptide of Group II.

The special technical feature of Group VI is a a method of identifying a compound that bind to the polypeptide of Group II.

The special technical feature of Group VII is a a method of treatment using the polypeptide of Group II.

The special technical feature of Group VIII is a a method of treatment using the antibody of Group III.

Accordingly, Groups I-X are not so linked by the same or a corresponding special technical feature as to form a single general inventive concept.